

PCSK9 ANTAGONIST COMPOUNDS

CROSS REFERENCE TO RELATED APPLICATION

This application claims priority to U.S. Application No. 62/687,913, filed June 21, 2018, which is incorporated herein by reference in its entirety.

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BACKGROUND OF THE INVENTION

The identification of compounds and/or agents that are effective in the treatment of cardiovascular affliction is highly desirable. In clinical trials, reductions in LDL cholesterol levels have been directly related to the rate of coronary events; Law *et al.*, 2003 *BMJ* 326:1423-1427. The moderate lifelong reduction in plasma LDL cholesterol levels was found to correlate with a substantial reduction in the incidence of coronary events; Cohen *et al.*, 2006 *N. Engl. J. Med.* 354:1264-1272. This was the case even in populations with a high prevalence of non-lipid-related cardiovascular risk factors; *supra*. Accordingly, there is great benefit to be reaped from the managed control of LDL cholesterol levels.

15 Proprotein convertase subtilisin-kexin type 9 (hereinafter called "PCSK9"), also known as neural apoptosis-regulated convertase 1 ("NARC-1"), is a proteinase K-like subtilase identified as the 9th member of the secretory subtilase family; see Seidah *et al.*, 2003 *PNAS* 100:928-933. PCSK9 belongs to the mammalian proprotein convertase family of serine proteases and contains an N-terminal signal sequence, a prodomain, a catalytic domain, and a C-terminal domain; see Seidah *et al.*, 2012 *Nat. Rev. Drug Discov.* 11:367–383. A study of PCSK9 transcriptional regulation demonstrated that it is regulated by sterol regulatory element-binding proteins ("SREBP"), as seen with other genes involved in cholesterol metabolism; Maxwell *et al.*, 2003 *J. Lipid Res.* 44:2109-2119, as is typical of other genes implicated in lipoprotein metabolism; Dubuc *et al.*, 2004 *Arterioscler. Thromb. Vasc. Biol.* 24:1454-1459. Statins have been shown to upregulate PCSK9 expression in a

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manner attributed to the cholesterol-lowering effects of the drugs; *supra*. Moreover, it has been shown that PCSK9 promoters possess two conserved sites involved in cholesterol regulation, a sterol regulatory element and an Sp1 site; *supra*.

While in the endoplasmic reticulum, PCSK9 performs as its only catalytic activity an autocleavage between residues Gln-152 and Ser-153; see Naureckiene *et al.*, 2003 *Arch. Biochem. Biophys.* 420:55–67; Seidah *et al.*, 2003 *Proc. Natl. Acad. Sci. U. S. A.* 100:928–933. The prodomain remains tightly associated with the catalytic domain during subsequent trafficking through the *trans*-Golgi network. The maturation via autocleavage has been demonstrated to be critical for PCSK9 secretion and subsequent extracellular function (see Benjannet *et al.*, 2012 *J. Biol. Chem.* 287:33745–33755). Accordingly, several lines of evidence demonstrate that PCSK9, in particular, lowers the amount of hepatic LDLR protein and thus compromises the liver's ability to remove LDL cholesterol from the circulation.

Adenovirus-mediated overexpression of PCSK9 in the liver of mice results in the accumulation of circulating LDL-C due to a dramatic loss of hepatic LDLR protein, with no effect on LDLR mRNA levels; Benjannet *et al.*, 2004 *J. Biol. Chem.* 279:48865–48875; Maxwell & Breslow, 2004 *PNAS* 101:7100–7105; Park *et al.*, 2004 *J. Biol. Chem.* 279:50630–50638; and Lalanne *et al.*, 2005 *J. Lipid Res.* 46:1312–1319. The effect of PCSK9 overexpression on raising circulating LDL-C levels in mice is completely dependent on the expression of LDLR, again, indicating that the regulation of LDL-C by PCSK9 is mediated through downregulation of LDLR protein. In agreement with these findings, mice lacking PCSK9 or in which PCSK9 mRNA has been lowered by antisense oligonucleotide inhibitors have higher levels of hepatic LDLR protein and a greater ability to clear circulating LDL-C; Rashid *et al.*, 2005 *PNAS* 102:5374–5379; and Graham *et al.*, 2007 *J. Lipid Res.* 48(4):763–767. In addition, lowering PCSK9 levels in cultured human hepatocytes by siRNA also results in higher LDLR protein levels and an increased ability to take up LDL-C;

Benjannet *et al.*, 2004 *J. Biol. Chem.* 279:48865-48875; and Lalanne *et al.*, 2005 *J. Lipid Res.* 46:1312-1319. Together, these data indicate that PCSK9 action leads to increased LDL-C by lowering LDLR protein levels.

A number of mutations in the gene PCSK9 have also been conclusively associated
5 **with autosomal dominant hypercholesterolemia ("ADH"), an inherited metabolism disorder characterized by marked elevations of low density lipoprotein ("LDL") particles in the**
plasma which can lead to premature cardiovascular failure; see Abifadel *et al.*, 2003 *Nature Genetics* 34:154-156; Timms *et al.*, 2004 *Hum. Genet.* 114:349-353; Leren, 2004 *Clin. Genet.* 65:419-422. A later-published study on the S127R mutation of Abifadel *et al.*,
10 *supra*, reported that patients carrying such a mutation exhibited higher total cholesterol and apoB100 in the plasma attributed to (1) an overproduction of apoB100-containing
lipoproteins, such as low density lipoprotein ("LDL"), very low density lipoprotein ("VLDL") and intermediate density lipoprotein ("IDL"), and (2) an associated reduction in clearance or
conversion of said lipoproteins; Ouguerram *et al.*, 2004 *Arterioscler. Thromb. Vasc. Biol.*
15 24:1448-1453.

Accordingly, there can be no doubt that PCSK9 plays a role in the regulation of LDL.
The expression or upregulation of PCSK9 is associated with increased plasma levels of
LDL cholesterol, and the corresponding inhibition or lack of expression of PCSK9 is
associated with reduced LDL cholesterol plasma levels. Decreased levels of LDL
20 cholesterol associated with sequence variations in PCSK9 have been found to confer
protection against coronary heart disease; Cohen, 2006 *N. Engl. J. Med.* 354:1264-1272.

Thus, identification of compounds and/or agents effective in the treatment of
cardiovascular affliction is highly desirable, including antagonism of PCSK9's role in LDL
regulation, however, in general, because PCSK9 circulates in blood and has modest
25 binding affinity to cell surface LDL receptors here-to-fore attempts to utilize this mechanism

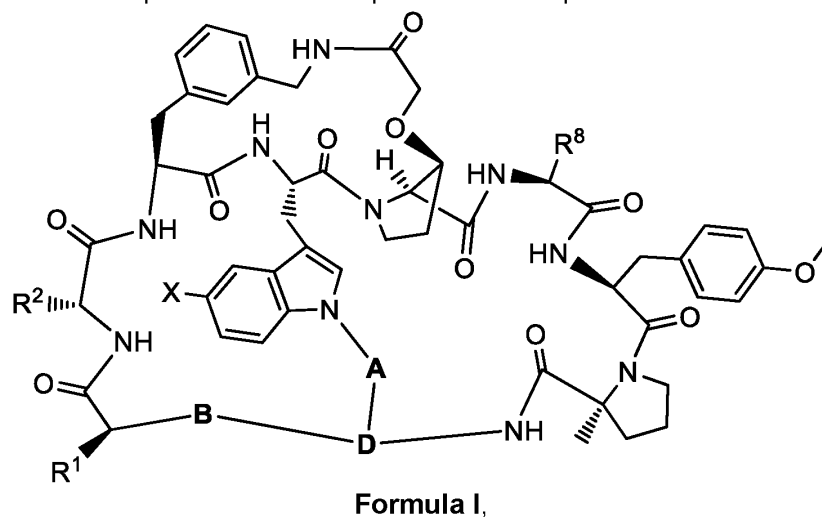
in treatment of diseases related to high serum LDL levels have been focused on the use of large biomolecules, for example, antibodies. Accordingly, there is scant publication reflecting activity toward this target using small peptides or small molecules to inhibit PCSK9, see for example, Zhang *et al.*, 2014 J. Biol. Chemistry, 289(2): 942-955.

Moreover, there is a paucity of compounds which are amenable to formulation into a dosage form for utilizing an oral administration route of dosing such compounds, a route which would be highly desirable for the provision of therapy for conditions in which regulation of the activities of PCSK9 could play a role.

The present invention advances these interests by providing antagonists of PCSK9 which are believed to be of use for inhibiting the activities of PCSK9 and the corresponding role PCSK9 plays in various conditions for which the administration of a PCSK9 antagonist provides therapy.

SUMMARY OF THE INVENTION

In one aspect the invention provides a compound of **Formula I**:

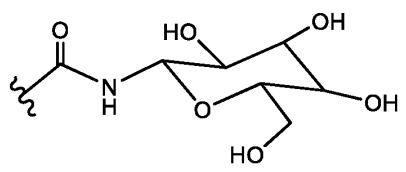


wherein:

X is H, F, Cl or Br;

R¹ is selected from:

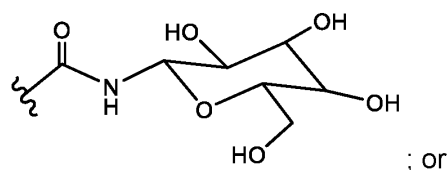
- (a) -H; or
- (b) $-(\text{CH}_2)_z\text{-R}^{14\text{A}}$, wherein: z is 1-6, and $\text{R}^{14\text{A}}$ is:
- (i) -H;
- (ii) $-\text{NH}_2$;
- (iii) $-\text{N}^+\text{H}_3$;
- (iv) $-\text{N}^+(\text{H}_3\text{C})_3$;
- (v) $-\text{NH-C(O)-}[(\text{CH}_2)_2\text{-O-}]_2\text{-(CH}_2)_2\text{R}^{14\text{B}}$ wherein $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
- (vi) $-\text{NH-C(O)-}[(\text{CH}_2)_{y12}\text{-O-}]_2\text{-(CH}_2)_{y13}\text{R}^{14\text{B}}$ wherein: y12 and y13 are not both 2 and are independently 2 to 4; and $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
- (vii) $-\text{NH-C(O)-}(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, y= 1 to 6 and $\text{R}^{14\text{C}}$ is $-\text{O-(CH}_2)_{z\text{a}}\text{-N}^+(\text{CH}_3)_3$, wherein $z\text{a}$ is 3 or 4; and
- (viii) $-\text{NH-C(O)-}(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, y= 1 to 6 and $\text{R}^{14\text{C}}$ is:
- (ai) $-\text{O-(CH}_2)_2\text{-N}^+(\text{CH}_3)_3$;
- (aii) $-\text{N}^+(\text{CH}_3)_3$; or
- (aiii) a moiety of the formula:



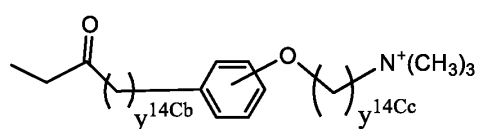
R^2 is selected from:

- (a) -H; and
- (b) $-(\text{CH}_2)_z\text{-R}^{14\text{A}}$, wherein: z is 1-6, and $\text{R}^{14\text{A}}$ is selected from:
- (i) -H;
- (ii) $-\text{NH}_2$;

- (iii) $-\text{N}^+\text{H}_3$;
- (iv) $-\text{N}^+(\text{H}_3\text{C})_3$;
- (v) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_2-\text{O}]_2-(\text{CH}_2)_2\text{R}^{14\text{B}}$ wherein $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
- 5 (vi) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_{y12}-\text{O}]_2-(\text{CH}_2)_{y13}\text{R}^{14\text{B}}$ wherein:
 $y12$ and $y13$ are not both 2 and are independently 2 to 4; and
 $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
- (vii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y=1$ to 6 and $\text{R}^{14\text{C}}$ is $-\text{O}-(\text{CH}_2)_{z\text{b}}-\text{N}^+(\text{CH}_3)_3$, wherein $z\text{b}$ is 3 or 4; and
- 10 (viii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y=1$ to 6 and $\text{R}^{14\text{C}}$ is:
- (ai) $-\text{O}-(\text{CH}_2)_2-\text{N}^+(\text{CH}_3)_3$;
- (aii) $-\text{N}^+(\text{CH}_3)_2\text{R}^{14\text{ca}}$, wherein $\text{R}^{14\text{ca}}$ is $-\text{CH}_3$ or $-(\text{CH}_2)_{1-4}-\text{OCH}_3$;
- (aiii) a moiety of the formula:

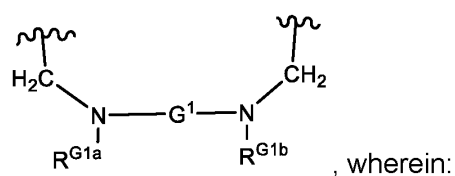


- 15 (aiv) a moiety of the formula:



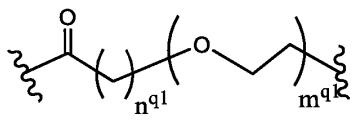
where $\text{R}^{14\text{Cb}}$ and $\text{R}^{14\text{Cc}}$ are 1 to 4; or

R^1 and R^2 may be bonded together to form a moiety of the formula:



G^1 , R^{G1a} and R^{G1b} are defined as follows:

(a) G^1 is a linker moiety of the formula:

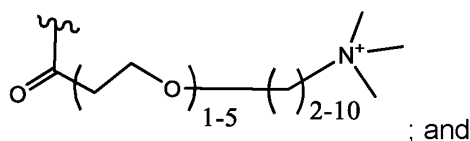


wherein n^{q1} is 1 to 6, m^{q1} is 0, 1 or 2 and together the value of n^{q1} and m^{q1} are selected such that the length of the linker moiety they define does not exceed a total of 8 carbon and/or oxygen atoms comprising the chain including the carbon atom in the chain that forms the carbonyl moiety;

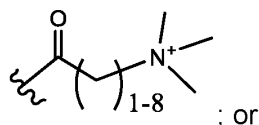
R^{G1a} is selected from: (i) $-H$; and (ii) alkyl of up to 4 carbon atoms; and

R^{G1b} is selected from:

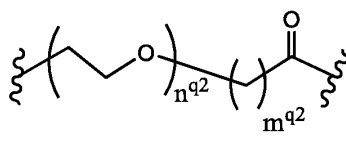
(i) a moiety of the formula:



(ii) a moiety of the formula:



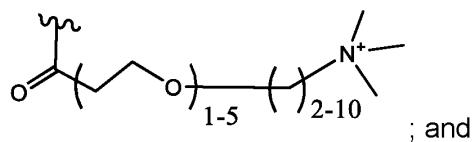
(b) G^1 is a linker moiety of the formula:



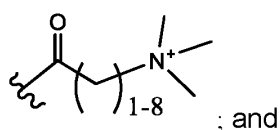
wherein n^{q2} is 0, 1 or 2, m^{q2} is 1 to 6, and together the value of n^{q2} and m^{q2} are selected such that the length of the linker moiety they define does not exceed a total of 8 carbon and/or oxygen atoms comprising the chain including the carbon atom in the chain that forms the carbonyl moiety;

R^{G1a} is selected from:

(i) a moiety of the formula:



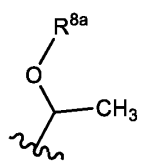
(ii) a moiety of the formula:



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R^{G1b} is selected from: (i) -H; and (ii) alkyl of up to 4 carbon atoms;

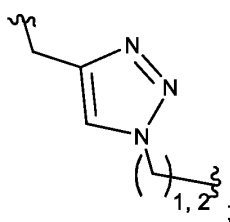
R⁸ is -CH₃ or a moiety of the formula:



wherein R^{8a} is -H, or a linear, branched or cyclic alkyl of up to four carbon atoms;

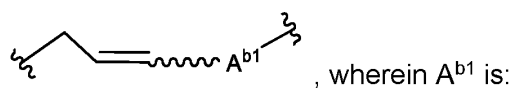
10 **A** is selected from:

(a) a moiety of the formula:



(b) -CH₂-(CH₂)_y-CH₂-, wherein y is 1 to 6;

(c) a moiety of the formula:

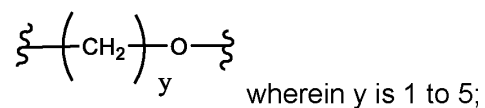


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(i) a moiety of the formula:



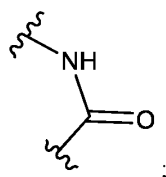
(ii) a moiety of the formula:



(d) a moiety of the formula: $-\text{CH}_2-(\text{CH}_2)_m-\text{O}-(\text{CH}_2)_n-$, wherein $m = 1 \text{ to } 5$, and $n = 0 \text{ or } 1 \text{ to } 4$;

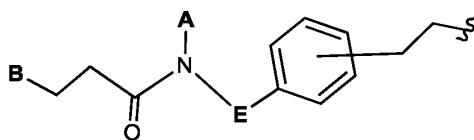
B is:

- (a) a bond;
- (b) $-(\text{CH}_2)_{1-4}$; or
- (c) a moiety of the formula:



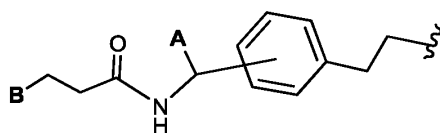
D is:

- (a) a moiety of the Formula:



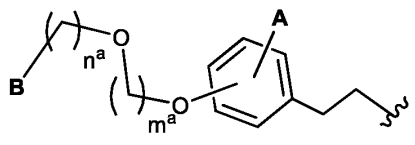
wherein **E** is $-\text{CH}_2-$ or $-(\text{CH}_2)_{2-4}-\text{O}-$, and **A** and **B** are as defined above;

- (b) a moiety of the formula:



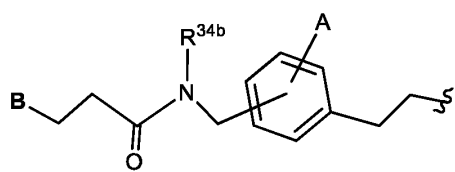
wherein **A** and **B** are as defined above;

(c) a moiety of the formula:



wherein n^a is 1, 2, or 3, m^a is 2, 3, or 4, and $n^a + m^a \geq 3$, and wherein **A** and **B** are as defined above;

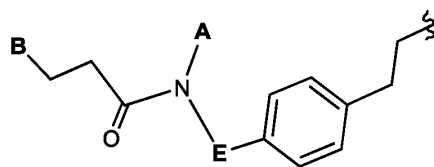
5 (d) a moiety of the formula:



wherein, R^{34b} is -H or a linear, branched or cyclic alkyl of up to four carbon atoms, and **A** and **B** are as defined above,

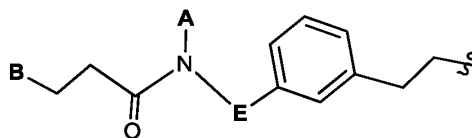
or a pharmaceutically acceptable salt of any thereof.

10 In a further embodiment, the invention provides a compound of Formula I, wherein **X** is F, or a pharmaceutically acceptable salt of any thereof. In some embodiments, it is preferred for **D** to be a moiety of the formula:



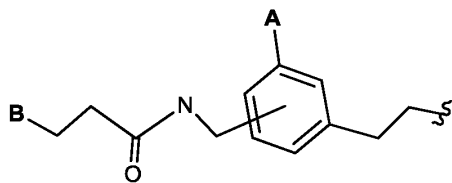
wherein, **E** is -CH₂- or -(CH₂)₂-O-, and **A** and **B** are as defined herein.

15 In some embodiments, it is preferred for **D** to be a moiety of the formula:



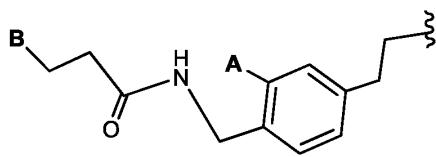
wherein, **E** is -CH₂- or -(CH₂)₂-O-, and **A** and **B** are as defined herein.

In some embodiments it is preferred for D to be a moiety of the Formula:



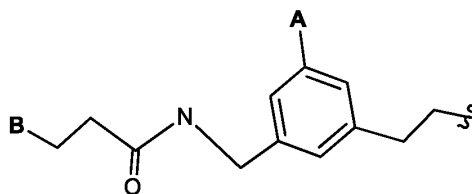
wherein **A** and **B** are as defined herein.

In some embodiments, it is preferred for **D** to be a moiety of the formula:



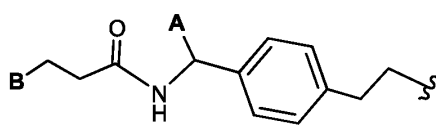
wherein **A** and **B** are as defined herein.

In some embodiments, it is preferred for **D** to be a moiety of the formula:



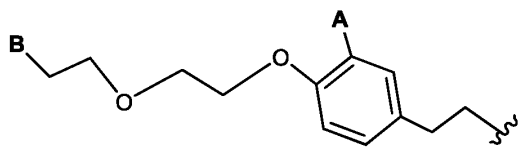
wherein **A** and **B** are as defined herein.

In some embodiments, it is preferred for **D** to be a moiety of the formula:



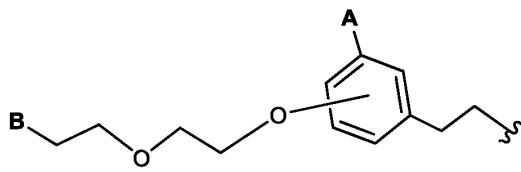
wherein **A** and **B** are as defined herein.

In some embodiments, it is preferred for **D** to be a moiety of the formula:



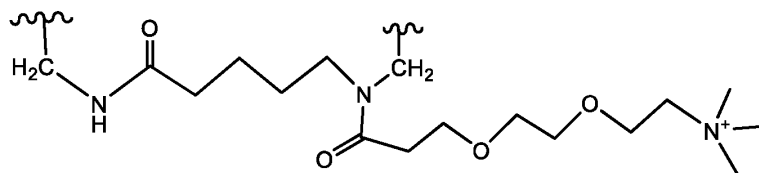
wherein **A** and **B** are as defined herein.

In some embodiments, it is preferred for **D** to be a moiety of the formula:



wherein **A** and **B** are as defined herein.

In some embodiments wherein R¹ and R² are joined together, along with the peptide
5 ring to which they are attached forming thereby a cyclic structure, it is preferred for R¹ and
R² to form a moiety of the structure:



In one embodiment the present invention provides pharmaceutical compositions
comprising a compound of the invention, for example, a compound of Formula I, and at
10 least one pharmaceutical excipient, preferably a composition directed to oral administration.

In one aspect the present invention provides a method of antagonizing PCSK9 in the
provision of therapy for disease states related to PCSK9 activity, for example,
atherosclerosis, hypercholesterolemia, coronary heart disease, metabolic syndrome, acute
coronary syndrome, or related cardiovascular disease and cardiometabolic conditions, by
15 administering to a subject in need thereof a therapeutically effective amount of a compound
of Formula I, or a salt thereof, preferably in the form of a pharmaceutical composition.

DETAILED DESCRIPTION OF THE INVENTION

In the description that follows conventional structural representation is employed and
20 includes conventional stereochemical notation for certain asymmetric carbon centers.

Thus, structural representation of compounds of the invention includes conventional stereochemical notation for some asymmetric carbon centers shown in the example compounds. Accordingly, in such instances, solid black “wedge” bonds represent bonds projecting from the plane of the reproduction medium, “hashed wedge” bonds representing descending bonds into the plane of the reproduction medium, and a “wavy” line appended to a carbon bearing a double bond indicates both possible cis and trans orientations are included. As is conventional, plain solid lines represent all spatial configurations for the depicted bonding. Accordingly, where no specific stereochemical notation is supplied, the representation contemplates all stereochemical and spatial orientations of the structural features.

As is shown in the examples of the invention, and mentioned above, particular asymmetric carbon centers are structurally represented using conventional “Solid Wedge” and “Hash Wedge” bonding representation. For the most part, absolute configuration has not been determined for the example compounds, but has been assigned by analogy to specific example compounds of known stereochemical configurations (determined by X-ray crystallography) prepared using the same or analogous reaction conditions and starting reagents and isolated under the same chromatographic conditions. Accordingly, specific assignment of the configurations structurally represented herein is meant to identify the specific compounds prepared having an excess of one particular stereoisomer and is not put forth herein necessarily as being a statement of the absolute determination of the stereochemical structure of said compound unless otherwise noted in the data presented.

It will be appreciated that where isomeric mixtures are obtained, the preparation of individual stereoisomers in significant percentages of enantiomeric excess can be carried out, if desired, by separation of the mixture using customary methods, for example by chromatography or crystallization, or by the use of stereochemically uniform starting

materials for the synthesis described, or by stereoselective synthesis. Optionally a derivatization can be carried out before a separation of stereoisomers. The separation of a mixture of stereoisomers can be carried out at an intermediate step during the synthesis of a compound of Formula I or it can be done on a final racemic product.

5 Where indicated herein, absolute stereochemistry is determined by X-ray crystallography of crystalline products or crystalline intermediates which are derivatized, if necessary, with a reagent containing a stereogenic center of known configuration. Unless a particular isomer, salt, solvate (including hydrates) or solvated salt of such racemate, enantiomer, or diastereomer is indicated, the present invention includes all such isomers,
10 as well as salts, solvates (including hydrates) and solvated salts of such racemates, enantiomers, diastereomers and mixtures thereof.

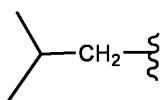
The present invention also embraces isotopically-labeled compounds of the present invention which are structurally identical to those recited herein, but for the fact that a statistically significant percentage of one or more atoms in that form of the compound are
15 replaced by an atom having an atomic mass or mass number different from the atomic mass or mass number of the most abundant isotope usually found in nature, thus altering the naturally occurring abundance of that isotope present in a compound of the invention. The present invention is meant to include all suitable isotopic variations of the compounds of Formula I.

20 Examples of isotopes that can be preferentially incorporated into compounds of the invention include isotopes of hydrogen, carbon, nitrogen, oxygen, phosphorus, iodine, fluorine and chlorine, for example, but not limited to: ^2H , ^3H , ^{11}C , ^{13}C , ^{14}C , ^{13}N , ^{15}N , ^{15}O , ^{17}O , ^{18}O , ^{31}P , ^{32}P , ^{35}S , ^{18}F , and ^{36}Cl , ^{123}I , and ^{125}I . It will be appreciated that other isotopes may be incorporated by known means also.

In particular, certain isotopically-labeled compounds of the invention (e.g., those labeled with ^3H , ^{11}C , and ^{14}C) are recognized as being particularly useful in compound and/or substrate tissue distribution assays using a variety of known techniques.

Additionally, compounds of the invention contemplate isotopic substitution include different isotopic forms of hydrogen (H), including protium (^1H) and deuterium (^2H or D). Protium is the predominant hydrogen isotope found in nature. Enriching for deuterium may afford certain therapeutic advantages, such as increasing *in vivo* half-life or reducing dosage requirements, or may provide a compound useful as a standard for characterization of biological samples. Isotopically-enriched compounds within Formula I can be prepared without undue experimentation by conventional techniques well known to those skilled in the art or by processes analogous to those described in the Schemes and Examples herein using appropriate isotopically-enriched reagents and/or intermediates.

Where a wavy line terminates a conventional bond (as opposed to connecting two atoms within a structure) it indicates a point of bonding to a structure, e.g.:



indicates a the secondary-butyl moiety is bonded via the methylene group via the bond terminated with the wavy line. Where an alphabetical notation is used to depict a substituent moiety, a dash is employed to indicate the point of bonding to the indicated substrate, e.g.: $-\text{CH}_2-\text{C}(\text{O})-\text{CH}_2\text{Cl}$ indicates the acetyl chloride moiety is bonded via the methylene portion of the moiety.

When any variable (e.g., n, R^a , R^b , etc.) occurs more than one time in any constituent or in Formula I, its definition on each occurrence is independent of its definition at every other occurrence unless otherwise specified at the point of definition. One of ordinary skill in the art will recognize that choice of combinations of the various substituents

defined in a structural representation, *i.e.* R^1 , R^A , etc., are to be chosen in conformity with well-known principles of chemical structure connectivity and stability, and combinations of substituents and/or variables are permissible only if such combinations result in stable compounds.

5 A "stable" compound is a compound which can be prepared and isolated and whose structure and properties remain or can be caused to remain essentially unchanged for a period of time sufficient to allow use of the compound for the purposes described herein (*e.g.*, therapeutic administration to a subject). The compounds of the present invention are limited to stable compounds embraced by Formula I.

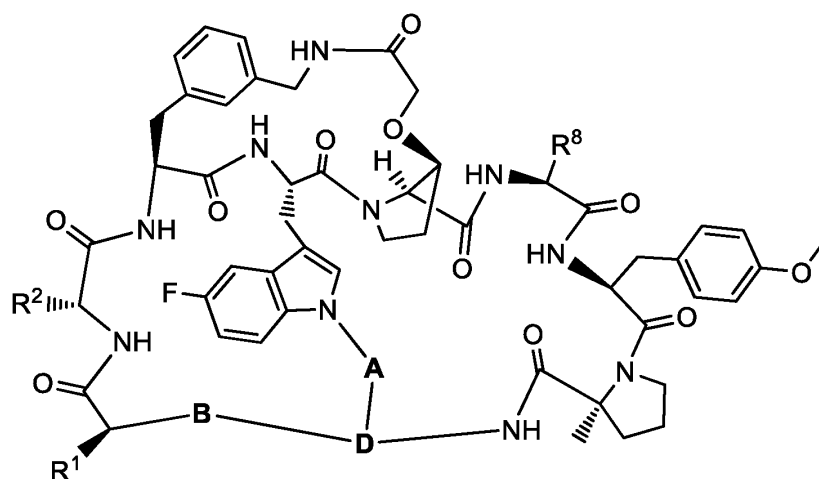
10 Where any variable or moiety is expressed in the form of a range, *e.g.* $(-CH_2-)_{1-4}$, both of the extremes of the specified range are included (*i.e.* 1 and 4 in the example) as well as all of the whole number values in between (*i.e.* 2 and 3 in the example).

The term "halogen" includes fluorine, chlorine, bromine and iodine unless specified otherwise at the point of use.

15 **As the term is used herein, "subjects" (alternatively "patients") refers to an animal,** preferably a mammal, and in particular a human or a non-human animal including livestock animals and domestic animals including, but not limited to, cattle, horses, sheep, swine, goats, rabbits, cats, dogs, and other mammals in need of treatment. In some embodiments the subject is preferably a human. As used herein, the term "administration" and variants thereof (*e.g.*, "administering" a compound) in reference to a compound of Formula I means
20 providing the compound, or a pharmaceutically acceptable salt thereof, to a subject in need of treatment.

As mentioned above, in one aspect the present invention includes the provision of compounds of **Formula I**, or a pharmaceutically acceptable salt thereof, which have properties that antagonize PCSK9 function.

In an embodiment, the compounds of Formula I have the structure of Formula IA:



Formula IA,

wherein:

R¹ is selected from:

(a) -H; or

(b) -(CH₂)_z-R^{14A}, wherein: z is 1-6, and R^{14A} is:

(i) -H;

(ii) -NH₂;

(iii) -N⁺H₃;

(iv) -N⁺(H₃C)₃;

(v) -NH-C(O)-[(CH₂)₂-O]₂-(CH₂)₂R^{14B} wherein R^{14B}

is: -NH₂; -N⁺H₃; -N(CH₃)₂; or -N⁺(CH₃)₃;

(vi) -NH-C(O)-[(CH₂)_{y12}-O]₁₋₄-(CH₂)_{y13}R^{14B}, preferably -NH-C(O)-[(CH₂)_{y12}-O]₂-(CH₂)_{y13}R^{14B} wherein:

y₁₂ and y₁₃ are not both 2 and are independently 2 to 4; and

R^{14B} is: -NH₂; -N⁺H₃; -N(CH₃)₂; or -N⁺(CH₃)₃;

(vii) -NH-C(O)-(CH₂)_yR^{14C}, wherein, y= 1 to 6 and R^{14C}

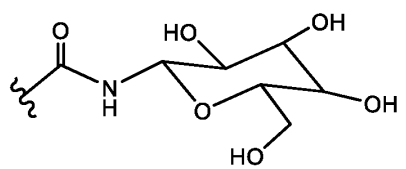
is -O-(CH₂)_{za}-N⁺(CH₃)₃, wherein za is 3 or 4; and

5 (viii) -NH-C(O)-(CH₂)_yR^{14C}, wherein, y= 1 to 6 and R^{14C} is:

(ai) -O-(CH₂)₂-N⁺(CH₃)₃;

(aii) -N⁺(CH₃)₃; or

(aiii) a moiety of the formula:



10 R² is selected from:

(a) -H; and

(b) -(CH₂)_z-R^{14A}, wherein: z is 1-6, and R^{14A} is selected from:

(i) -H;

(ii) -NH₂;

15 (iii) -N⁺H₃;

(iv) -N⁺(H₃C)₃;

(v) -NH-C(O)-[(CH₂)₂-O]₁₋₄-(CH₂)₂R^{14B}, preferably -NH-C(O)-[(CH₂)₂-O]₂-
(CH₂)₂R^{14B} wherein R^{14B} is: -NH₂; -N⁺H₃; -N(CH₃)₂; or -N⁺(CH₃)₃;

(vi) -NH-C(O)-[(CH₂)_{y₁₂}-O]₂-(CH₂)_{y₁₃}R^{14B} wherein:

20 y₁₂ and y₁₃ are not both 2 and are independently 2 to 4; and

R^{14B} is: -NH₂; -N⁺H₃; -N(CH₃)₂; or -N⁺(CH₃)₃;

(vii) -NH-C(O)-(CH₂)_yR^{14C}, wherein, y= 1 to 6 and R^{14C}

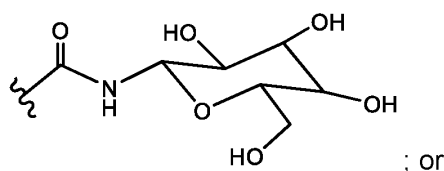
is -O-(CH₂)_{zb}-N⁺(CH₃)₃, wherein zb is 3 or 4; and

(viii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y=1$ to 6 and $\text{R}^{14\text{C}}$ is:

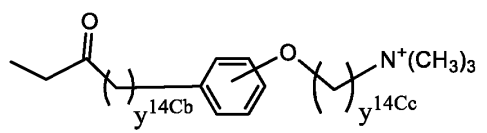
(ai) $-\text{O}-(\text{CH}_2)_2-\text{N}^+(\text{CH}_3)_3$;

(aii) $-\text{N}^+(\text{CH}_3)_2\text{R}^{14\text{ca}}$, wherein $\text{R}^{14\text{ca}}$ is $-\text{CH}_3$ or $-(\text{CH}_2)_{1-4}-\text{OCH}_3$;

(aiii) a moiety of the formula:

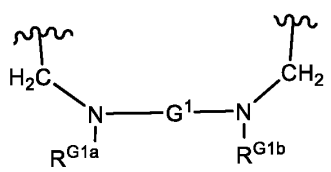


(aiv) a moiety of the formula:



where $\text{Y}^{14\text{Cb}}$ and $\text{Y}^{14\text{Cc}}$ are 1 to 4 ; or

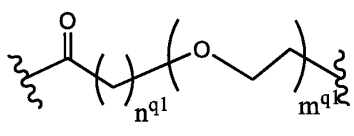
R^1 and R^2 may be bonded together to form a moiety of the formula:



wherein:

G^1 , $\text{R}^{\text{G}1\text{a}}$ and $\text{R}^{\text{G}1\text{b}}$ are defined as follows:

(a) G^1 is a linker moiety of the formula:

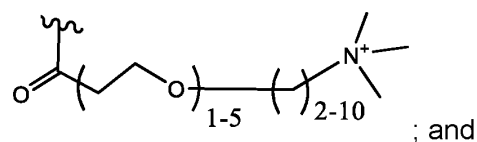


wherein $n^{\text{q}1}$ is 1 to 6 , $m^{\text{q}1}$ is 0 , 1 or 2 and together the value of $n^{\text{q}1}$ and $m^{\text{q}1}$ are selected such that the length of the linker moiety they define does not exceed a total of 8 carbon and/or oxygen atoms comprising the chain including the carbon atom in the chain that forms the carbonyl moiety;

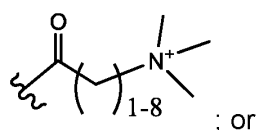
R^{G1a} is selected from: (i) -H; and (ii) alkyl of up to 4 carbon atoms; and

R^{G1b} is selected from:

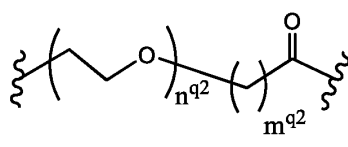
(i) a moiety of the formula:



(ii) a moiety of the formula:



(b) G¹ is a linker moiety of the formula:



wherein n^{q2} is 0, 1 or 2, m^{q2} is 1 to 6, and together the value of n^{q2} and m^{q2}

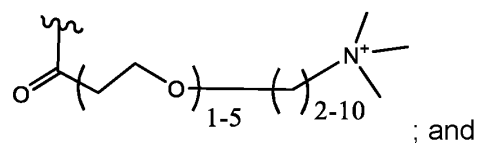
are selected such that the length of the linker moiety they define does not

exceed a total of 8 carbon and/or oxygen atoms comprising the chain

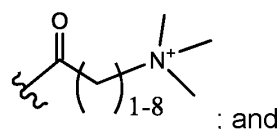
including the carbon atom in the chain that forms the carbonyl moiety;

R^{G1a} is selected from:

(i) a moiety of the formula:

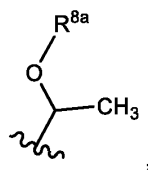


(ii) a moiety of the formula:



R^{G1b} is selected from: (i) -H; and (ii) alkyl of up to 4 carbon atoms;

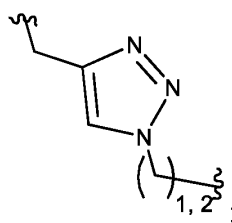
R⁸ is -CH₃ or a moiety of the formula:



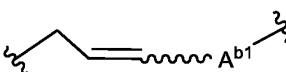
wherein R^{8a} is -H, or a linear, branched or cyclic alkyl of up to four carbon atoms;

A is selected from:

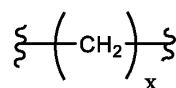
- 5 (a) a moiety of the formula:



- (b) -CH₂-(CH₂)_y-CH₂-, wherein y is 1 to 6;

- (c) a moiety of the formula: , wherein A^{b1} is:

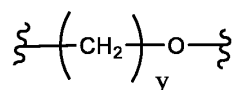
- (i) a moiety of the formula:



10

wherein x is 1 to 6; or

- (ii) a moiety of the formula:



wherein y is 1 to 5;

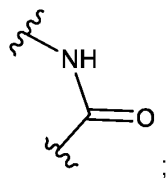
- 15 (d) a moiety of the formula: -CH₂-(CH₂)_m-O-(CH₂)_n-, wherein m = 1 to 5, and n = 0 or 1 to 4;

B is:

- (a) a bond;

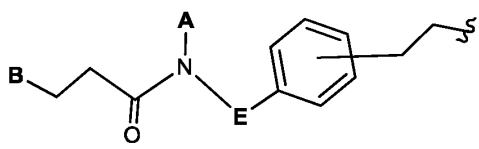
(b) $-(CH_2)_{1-4}$; or

(c) a moiety of the formula:



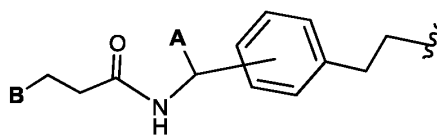
D is:

5 (a) a moiety of the Formula:



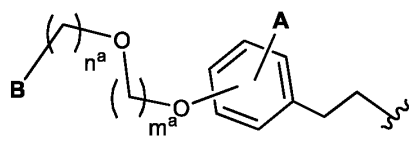
wherein **E** is $-CH_2-$ or $-(CH_2)_{2-4}-O-$, and **A** and **B** are as defined above;

(b) a moiety of the formula:



wherein **A** and **B** are as defined above;

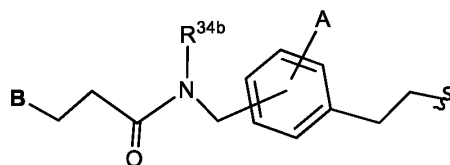
(c) a moiety of the formula:



wherein n^a is 1, 2, or 3, m^a is 2, 3, or 4, and $n^a + m^a$ is at least 3, and wherein **A** and **B** are

15 as defined above;

(d) a moiety of the formula:



wherein, R^{34b} is $-H$ or a linear, branched or cyclic alkyl of up to four carbon atoms, and **A** and **B** are as defined above,

or a pharmaceutically acceptable salt of any thereof.

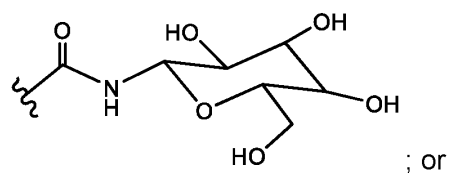
5 In an embodiment of the compounds of Formula IA, R^1 is $-(CH_2)_z-R^{14A}$, wherein: z is 1-6, and R^{14A} is:

- (i) $-H$;
- (ii) $-NH_2$;
- (iii) $-N^+H_3$; or
- 10 (iv) $-N^+(H_3C)_3$;

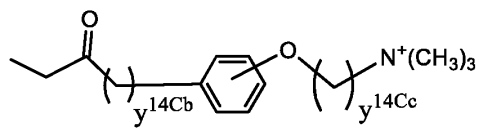
R^2 is $-(CH_2)_z-R^{14A}$, wherein: z is 1-6, and R^{14A} is selected from:

- (i) $-H$;
- (ii) $-NH_2$;
- (iii) $-NH-C(O)-[(CH_2)_2-O]_{1-4}-(CH_2)_2R^{14B}$, preferably $-NH-C(O)-[(CH_2)_2-O]_2-(CH_2)_2R^{14B}$ wherein R^{14B} is: $-NH_2$; $-N^+H_3$; $-N(CH_3)_2$; or $-N^+(CH_3)_3$;
- 15 (iv) $-NH-C(O)-[(CH_2)_{y12}-O]_2-(CH_2)_{y13}R^{14B}$ wherein:
 y_{12} and y_{13} are not both 2 and are independently 2 to 4; and
 R^{14B} is: $-NH_2$; $-N^+H_3$; $-N(CH_3)_2$; or $-N^+(CH_3)_3$;
- (v) $-NH-C(O)-(CH_2)_yR^{14C}$, wherein, $y = 1$ to 6 and R^{14C}
20 is $-O-(CH_2)_{zb}-N^+(CH_3)_3$, wherein zb is 3 or 4; and
- (vi) $-NH-C(O)-(CH_2)_yR^{14C}$, wherein, $y = 1$ to 6 and R^{14C} is:
 - (ai) $-O-(CH_2)_2-N^+(CH_3)_3$;
 - (aii) $-N^+(CH_3)_2R^{14ca}$, wherein R^{14ca} is $-CH_3$ or $-(CH_2)_{1-4}-OCH_3$;

(aiii) a moiety of the formula:



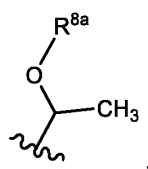
(aiv) a moiety of the formula:



5

where Y^{14Cb} and Y^{14Cc} are 1 to 4; or

R^8 is $-CH_3$ or a moiety of the formula:

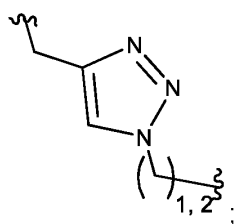


wherein R^{8a} is $-H$, or a linear, branched or cyclic alkyl of up to four carbon atoms;

A is selected from:

10

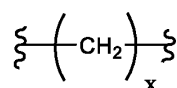
(a) a moiety of the formula:



(b) $-CH_2-(CH_2)_y-CH_2-$, wherein y is 1 to 6;

(c) a moiety of the formula: , wherein A^{b1} is:

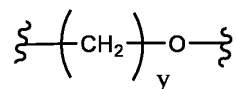
(i) a moiety of the formula:



15

wherein x is 1 to 6; or

(ii) a moiety of the formula:



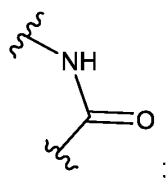
wherein y is 1 to 5; and

5 (d) a moiety of the formula: $-\text{CH}_2-(\text{CH}_2)_m-\text{O}-(\text{CH}_2)_n-$, wherein m = 1 to 5, and n = 0 or 1 to 4;

B is:

(a) $-(\text{CH}_2)_{1-4}$; or

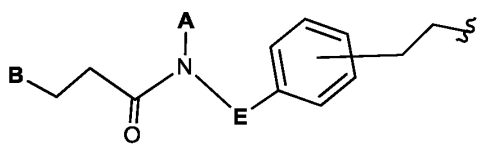
(b) a moiety of the formula:



10

D is:

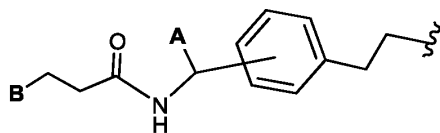
(a) a moiety of the Formula:



15

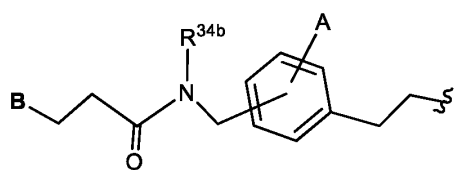
wherein **E** is $-\text{CH}_2-$ or $-(\text{CH}_2)_{2-4}-\text{O}-$, and **A** and **B** are as defined above;

(b) a moiety of the formula:



wherein **A** and **B** are as defined above; or

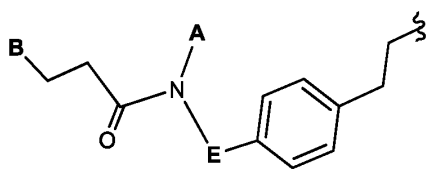
(c) a moiety of the formula:



wherein, R^{34b} is $-H$ or a linear, branched or cyclic alkyl of up to four carbon atoms,
and **A** and **B** are as defined above,

or a pharmaceutically acceptable salt of any thereof.

5 In an embodiment of the compound of Formula IA, **D** is a moiety of the formula:

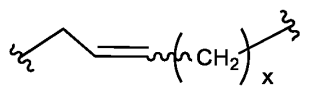


wherein, **E** is $-\text{CH}_2-$ or $-(\text{CH}_2)_2\text{O}-$, and **A** and **B** are as defined above in Formula IA.

In an embodiment of the compound of Formula IA, **A** is:

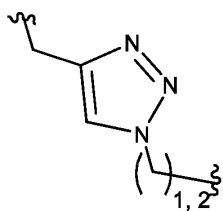
(a) $-(\text{CH}_2)_6$;

10 (b) a moiety of the formula:



wherein x is 1 to 3; or

(c) a moiety of the formula:

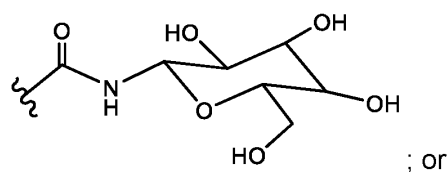


15 In another embodiment of the compound of Formula IA, R^2 is:

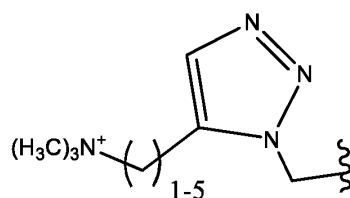
(a) $-(\text{CH}_2)_z\text{R}^{14A}$, wherein: z is 1-6, and R^{14A} is:

(a) $-H$;

- (b) $-\text{CH}_3$;
- (c) $-\text{NH}_2$;
- (d) $-\text{N}^+\text{H}_3$;
- (e) $-\text{N}^+(\text{H}_3\text{C})_3$;
- 5 (f) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_{2-4}-\text{O}]_{2-4}-(\text{CH}_2)_{2-4}\text{R}^{14\text{B}}$ wherein $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
- (g) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_y\text{R}^{14\text{C}}]$, wherein, $y = 1$ to 6 and $\text{R}^{14\text{C}}$ is:
- (ai) $-\text{O}-(\text{CH}_2)_{2-4}-\text{N}^+(\text{CH}_3)_3$;
- (aii) $-\text{N}^+(\text{CH}_3)_3$; or
- 10 (aiii) a moiety of the formula:



- (b) a moiety of the formula

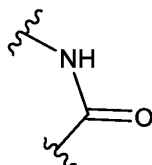


In a further embodiment of the compound of Formula IA, R^1 is selected from:

- 15 (a) $-\text{H}$;
- (b) $-(\text{CH}_2)_z-\text{R}^{14\text{A}}$, wherein: z is $1-6$, and $\text{R}^{14\text{A}}$ is:
- (i) $-\text{H}$;
- (ii) $-\text{N}^+\text{H}_3$; or
- (iii) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_2-\text{O}]_2-(\text{CH}_2)_2-\text{N}^+(\text{CH}_3)_3$.

In yet another embodiment of the compound of Formula IA, **A** is $-\text{CH}_2-(\text{CH}_2)_y-\text{CH}_2-$, wherein y is 3-5. In a further embodiment, **A** is $-(\text{CH}_2)_6$.

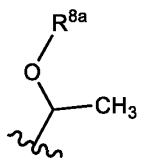
In another embodiment of the compound of Formula IA, **B** is a moiety of the formula:



5 In another embodiment of the compound of Formula IA, R^1 is $-(\text{CH}_2)_z-\text{R}^{14\text{A}}$, wherein: z is 1-6, and $\text{R}^{14\text{A}}$ is $-\text{H}$. In another embodiment of the compound of Formula IA, R^1 is $-(\text{CH}_2)_z-\text{R}^{14\text{A}}$, wherein: z is 1, and $\text{R}^{14\text{A}}$ is $-\text{H}$.

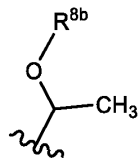
In another embodiment of the compound of Formula IA, R^2 is $-(\text{CH}_2)_z-\text{R}^{14\text{A}}$, wherein: z is 1-6, and $\text{R}^{14\text{A}}$ is $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein $y = 1$ to 6 and $\text{R}^{14\text{C}}$ is $-\text{N}^+(\text{CH}_3)_2\text{R}^{14\text{ca}}$,
 10 wherein $\text{R}^{14\text{ca}}$ is $-\text{CH}_3$.

In another embodiment of the compound of Formula IA, R^8 is a moiety of the formula:



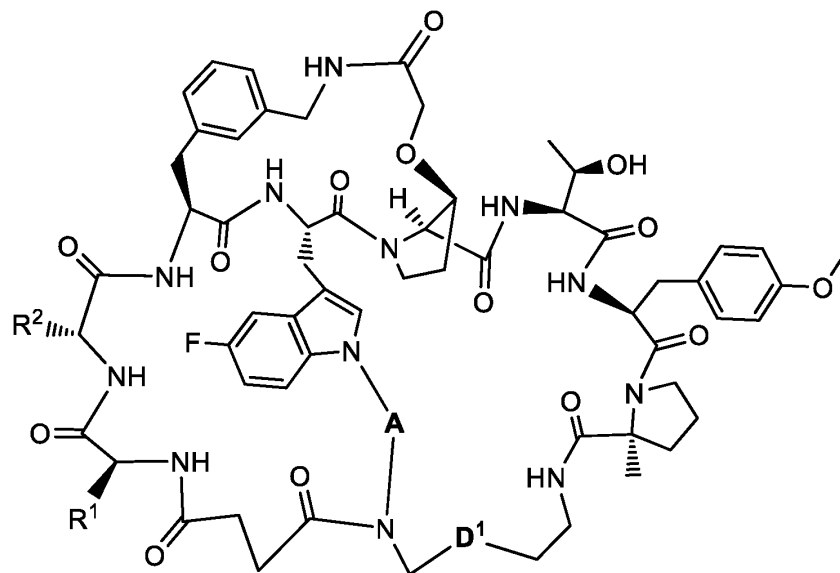
wherein $\text{R}^{8\text{a}}$ is $-\text{H}$ or linear alkyl of up to four carbon atoms. In a further embodiment, R^8 is

15 a moiety of the formula:

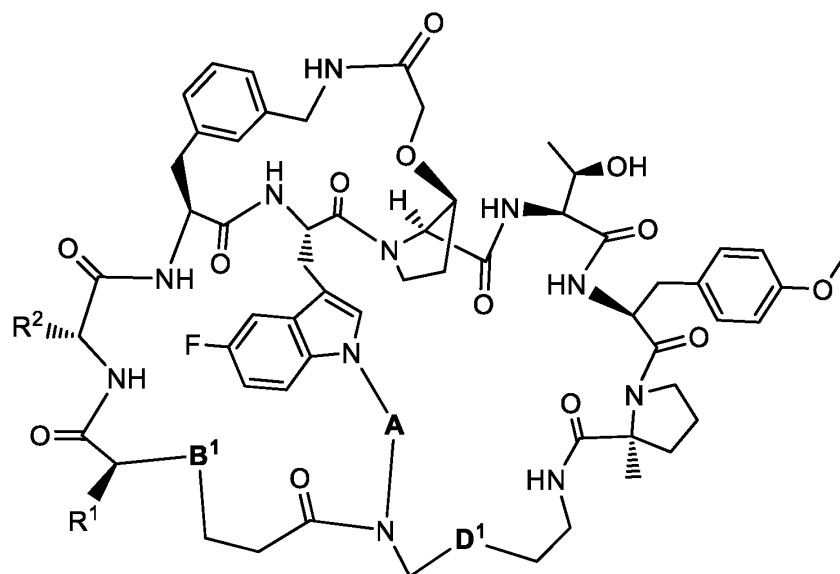


wherein $\text{R}^{8\text{b}}$ is $-\text{H}$, $-\text{CH}_3$, or $-\text{C}(\text{CH}_3)_3$.

In some embodiments, it is preferred for the compounds of **Formula I** to have the structure of **Formula II** or **Formula IIA**, or a pharmaceutically acceptable salt thereof:



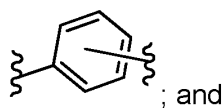
Formula II,



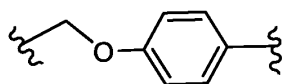
Formula IIA,

wherein, **A**, **R**¹ and **R**² are as defined above in Formula IA and **B**¹ is $-(CH_2)_{0-2}$, and **D**¹ is selected from:

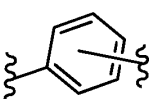
a) a moiety of the formula:

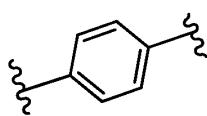


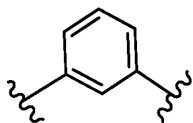
b) a moiety of the formula:



5 In some embodiments of **Formula II** or **Formula IIA**, it is preferred for **D¹** to be a

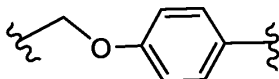
moiety of the formula: . In some embodiments **Formula II** or **Formula IIA**, it is

preferred for **D¹** to be a moiety of the formula:  In some embodiments of **Formula II** or **Formula IIA**, it is preferred for **D¹** to be a moiety of the formula:

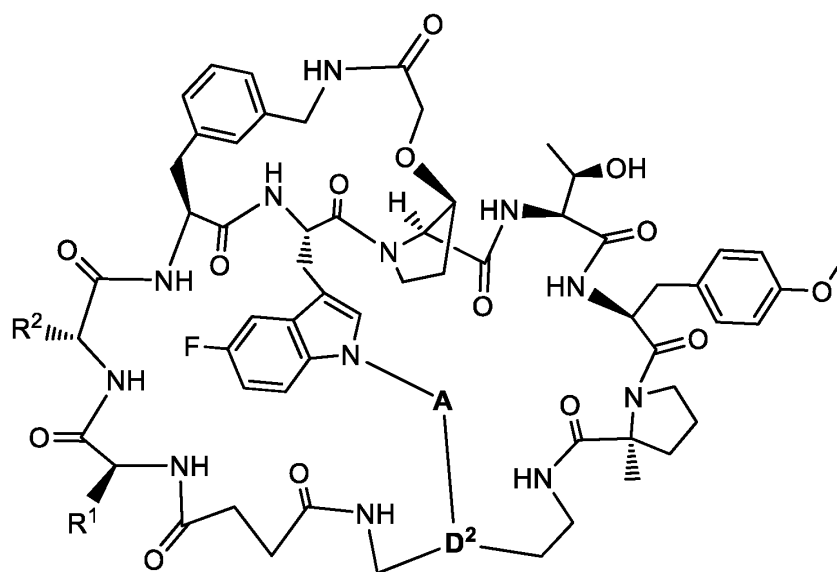


In some embodiments of **Formula II** or **Formula IIA**, it is preferred for **D¹**

10 to be a moiety of the formula:

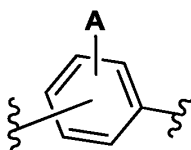


In some embodiments, the compound of **Formula I** is preferably a compound of **Formula III**:



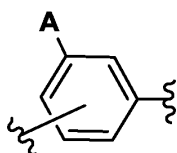
Formula III,

wherein, **A**, R^1 and R^2 are as defined above in Formula IA and D^2 is a moiety of the formula:

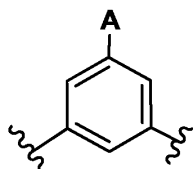


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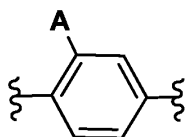
In some embodiments of **Formula III**, it is preferred for D^2 to be a moiety of the



formula: In some embodiments of **Formula III**, it is preferred for D^2 to be a



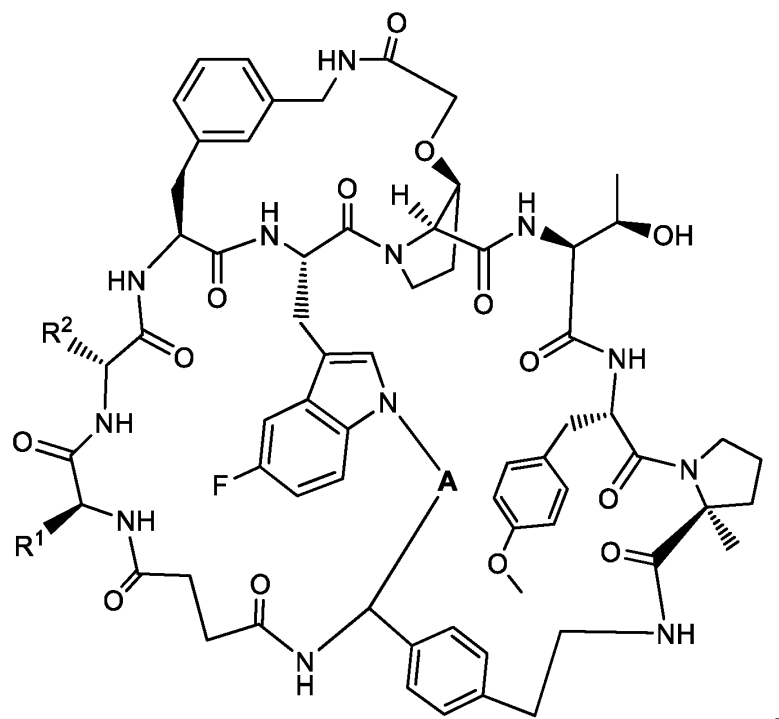
moiety of the formula: In some embodiments of **Formula III**, it is preferred



for D^2 to be a moiety of the formula:

In some embodiments, the compound of **Formula I** is preferably a compound of

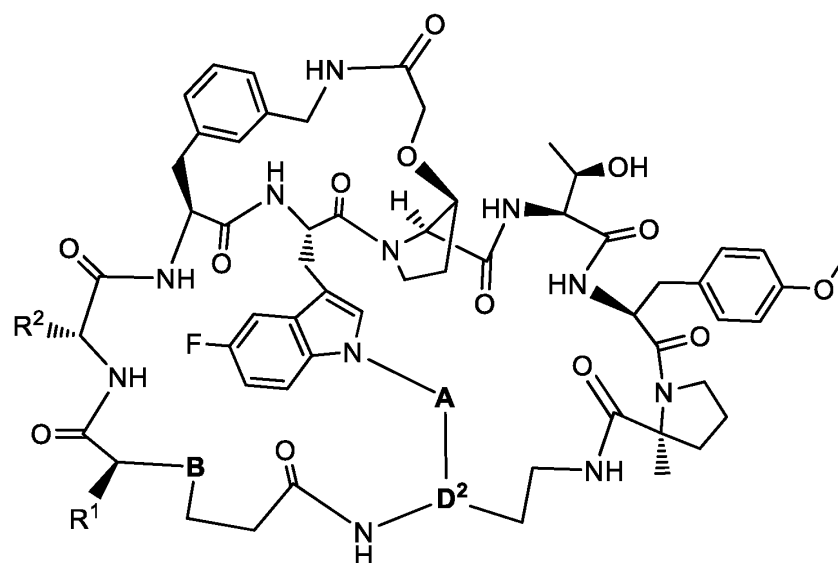
Formula IV:



Formula IV,

5 wherein, **A**, R¹ and R² are as defined above in Formula IA.

In some embodiments the compound of Formula I is a compound of Formula V:



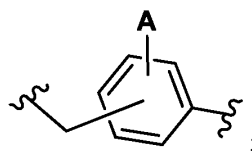
Formula V,

wherein

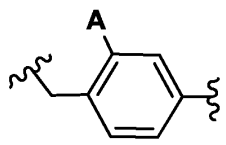
A, **B**, **R**¹ and **R**² are as defined above in Formula IA; and

D² is:

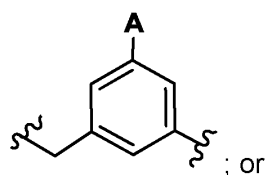
(a) a moiety of the formula:



(b) a moiety of the formula:

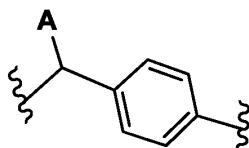


(c) a moiety of the formula:



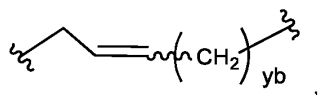
; or

(d) a moiety of the formula:



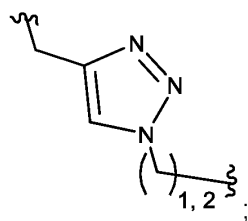
In some embodiments of **Formula I**, **Formula IA**, **Formula II**, **Formula IIA**, **Formula**

5 **III**, **Formula IV**, or **Formula V**, it is preferred for **A** to be a moiety of the formula: $-(CH_2)_{y_a}$, wherein y_a is 4 to 6. In some embodiments of **Formula I**, **Formula IA**, **Formula II**, or **Formula IIA**, it is preferred for **A** to be a moiety of the formula: $-CH_2-(CH_2)_{m_a}-O-(CH_2)_{n_a}-$, wherein m_a is 2 or 3 and n_a is 0 or 1. In some embodiments of **Formula III**, **Formula IV**, or **Formula V**, it is preferred for **A** to be a moiety of the formula: $-CH_2-(CH_2)_{m_a}-O-(CH_2)_{n_a}-$,
 10 wherein m_a is 2 or 4 and n_a is 0, 1, or 2. In some embodiments of **Formula I**, **Formula IA**, **Formula II**, **Formula IIA**, **Formula III**, **Formula IV**, or **Formula V**, it is preferred for **A** to be a moiety of the formula:



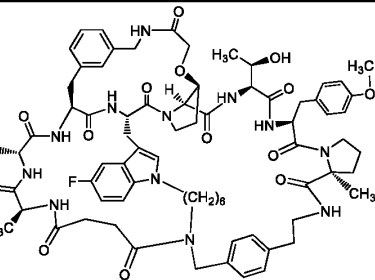
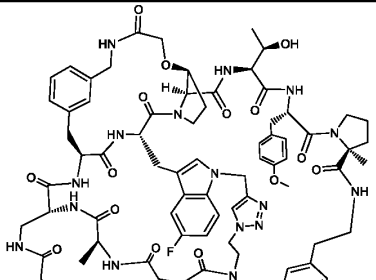
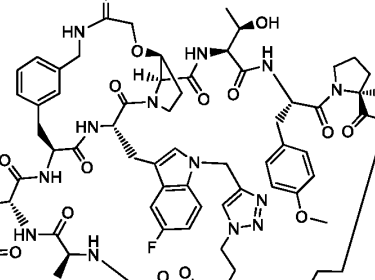
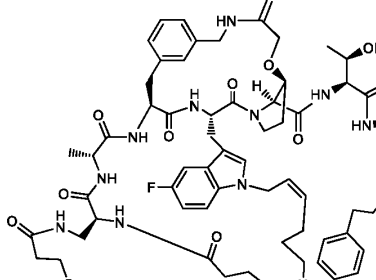
wherein y_b is 1 to 3. In some embodiments of **Formula I**, **Formula IA**, **Formula II**,

15 **Formula IIA**, **Formula III**, **Formula IV**, or **Formula V**, it is preferred for **A** to be a moiety of the formula:



Also provided herein as compounds of Formula I are compounds Ex-1, Ex-2, Ex-3, Ex-4, Ex-5, Ex-6, Ex-7, Ex-8, Ex-9, Ex-10, Ex-11, Ex-12, Ex-13, Ex-14, Ex-15, Ex-16, Ex-17, Ex-18, Ex-19, Ex-20, Ex-21, Ex-22, Ex-23, Ex-24, Ex-25, Ex-26, Ex-27, Ex-28, Ex-29, Ex-31, Ex-35, Ex-36, Ex-38, Ex-39, Ex-40, Ex-41, Ex-44, Ex-47, Ex-48, Ex-49, Ex-50, Ex-51, Ex-52, Ex-53, Ex-54, Ex-55, Ex-56, Ex-57, Ex-58, Ex-59, Ex-60, and Ex-61, or any pharmaceutically acceptable salt thereof. These compounds, which are disclosed in Table 1, are also referred to herein as “compounds of the invention.”

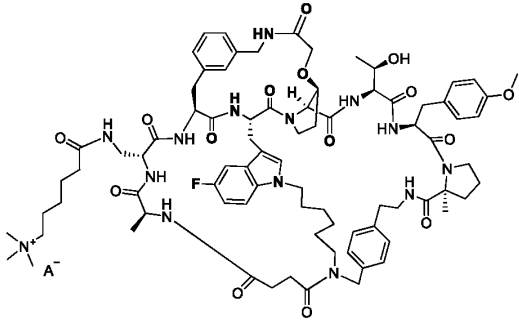
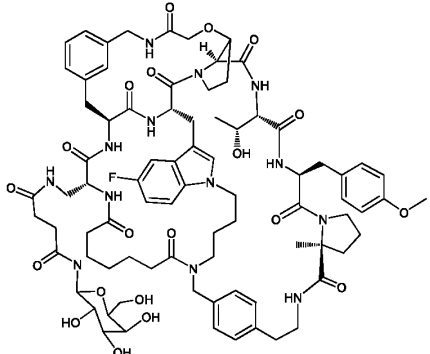
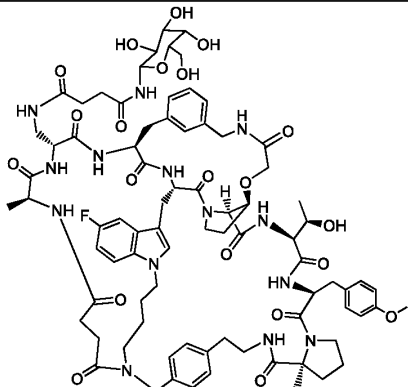
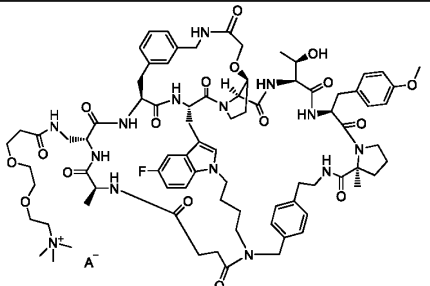
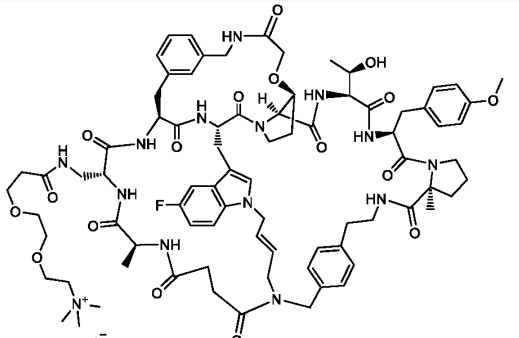
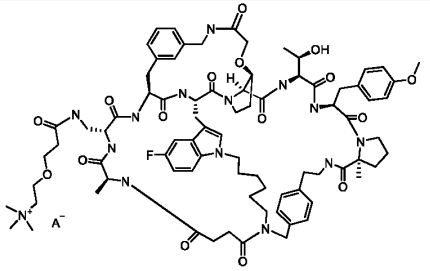
Table 1

Ex No	Structure	Ex No	Structure
Ex-01		Ex-02	
Ex-03		Ex-04	

Ex-05		Ex-06	
Ex-07		Ex-08	
Ex-09		Ex-10	
Ex-11		Ex-12	

Ex No	Structure	Ex No	Structure
Ex-13		Ex-14	
Ex-15		Ex-16	
Ex-17		Ex-18	

Ex No	Structure	Ex No	Structure
Ex-19		Ex-20	
Ex-21		Ex-22	
Ex-23		Ex-24	

Ex No	Structure	Ex No	Structure
Ex-25		Ex-26	
Ex-27		Ex-28	
Ex-29		Ex-31	

Ex No	Structure	Ex No	Structure
Ex-35		Ex-36	
Ex-38		Ex-39	
Ex-40		Ex-41	

5

10

15

Ex No	Structure	Ex No	Structure
Ex-44		Ex-47	
Ex-48		Ex-49	
Ex-50		Ex-51	

Ex No	Structure	Ex No	Structure
Ex-52		Ex-53	
Ex-54		Ex-55	
Ex-56		Ex-57	

Ex No	Structure	Ex No	Structure
Ex-58		Ex-59	
Ex-60		Ex-61	

wherein A⁻ is a pharmaceutically acceptable anion.

The term “salt(s)”, and its use in the phrase “pharmaceutically acceptable salts”

employed herein, includes any of the following: acidic salts formed with inorganic and/or organic acids, basic salts formed with inorganic and/or organic bases, zwitterionic and quaternary ammonium complexes. Salts of compounds of the invention may be formed by methods known to those of ordinary skill in the art, for example, by reacting a compound of the invention with an amount of acid or base, such as an equivalent amount, in a medium such as one in which the salt precipitates or in aqueous medium followed by lyophilization.

Compounds of the invention contain tri-coordinate nitrogen atoms, for example, primary, secondary or tertiary amino moieties, wherein, as is known, the lone pair of electrons residing on the nitrogen atom may be protonated with an appropriate acid or alkylated with an appropriate reagent, for example, alkyl bromide, under the appropriate

reaction conditions to provide tetracoordinate charged nitrogen stabilized by an anion generated in the process, for example, a halogen ion or conjugate base. Accordingly, compounds of the invention may be prepared in the form of a free-base or isolated in the form of a quaternary complex or a salt complex. In some instances where there is an appropriate acidic proton proximal to a basic nitrogen formation of a zwitterionic complex is possible. As the term is employed herein, salts of the inventive compounds, whether acidic salts formed with inorganic and/or organic acids, basic salts formed with inorganic and/or organic bases, salts formed which include zwitterionic character, for example, where a compound contains both a basic moiety, for example, but not limited to, a nitrogen atom, for example, an amine, pyridine or imidazole, and an acidic moiety, for example, but not limited to a carboxylic acid, and quaternary ammonium complexes are included in the scope of the inventive compounds described herein.

Accordingly, structural representation of compounds of the invention, whether in a free-base form, a salt form, a zwitterionic form or a quaternary ammonium form, also include all other forms of such compounds discussed above. Thus, one aspect of the invention is the provision of compounds of the invention in the form of a pharmaceutically acceptable salt, zwitterionic complex or quaternary ammonium complex. Those skilled in the art will recognize those instances in which the compounds of the invention may form such complexes, including where a tetracoordinate nitrogen can be quaternized or protonated and the charged nitrogen form stabilized by an associated anion. The term "pharmaceutically acceptable salt" refers to a salt (including a quaternary ammonium complex and an inner salt such as a zwitterion complex) which possesses effectiveness similar to or greater than a free-base form of the compound and which is not biologically or otherwise undesirable (e.g., is neither toxic nor otherwise deleterious to the recipient thereof).

The formation of pharmaceutically useful salts from basic (or acidic) pharmaceutical compounds are discussed, for example, by S. Berge et al., Journal of Pharmaceutical Sciences (1977) 66(1) 1-19; P. Gould, International J. of Pharmaceutics (1986) 33 201-217; Anderson et al, The Practice of Medicinal Chemistry (1996), Academic Press, New York; in
5 The Orange Book (Food & Drug Administration, Washington, D.C. on their website); and P. Heinrich Stahl, Camille G. Wermuth (Eds.), Handbook of Pharmaceutical Salts: Properties, Selection, and Use, (2002) Int'l. Union of Pure and Applied Chemistry, pp. 330-331. These disclosures are incorporated herein by reference.

The present invention contemplates both freebase forms of the compounds of the
10 invention and all available salts, including salts which are generally recognized as safe for use in preparing pharmaceutical formulations and those which may be formed presently **within the ordinary skill in the art and are later classified as being "generally recognized as safe" for use in the preparation of pharmaceutical formulations, termed herein as "pharmaceutically acceptable salts."** As will be appreciated, freebase compounds may be
15 prepared by controlling the conditions of isolation of the compound during synthesis or by neutralization and ion exchange from salt forms of compounds of the invention.

Examples of pharmaceutically acceptable acid salts include, but are not limited to, acetates, including trifluoroacetate salts, adipates, alginates, ascorbates, aspartates, benzoates, benzenesulfonates, bisulfates, borates, butyrates, citrates, camphorates,
20 camphorsulfonates, cyclopentanepropionates, digluconates, dodecylsulfates, ethanesulfonates, fumarates, glucoheptanoates, glycerophosphates, hemisulfates, heptanoates, hexanoates, hydrochlorides, hydrobromides, hydroiodides, 2-hydroxyethanesulfonates, lactates, maleates, methanesulfonates, methyl sulfates, 2-naphthalenesulfonates, nicotines, nitrates, oxalates, pamoates, pectinates, persulfates, 3-
25 phenylpropionates, phosphates, picrates, pivalates, propionates, salicylates, succinates,

sulfates, sulfonates (such as those mentioned herein), tartarates, thiocyanates, toluenesulfonates (also known as tosylates,) undecanoates, and the like.

Examples of pharmaceutically acceptable basic salts include, but are not limited to, ammonium salts, alkali metal salts such as sodium, lithium, and potassium salts, alkaline earth metal salts such as calcium and magnesium salts, aluminum salts, zinc salts, salts with organic bases (for example, organic amines) such as benzathines, diethylamine, dicyclohexylamines, hydrabamines (formed with N,N-bis(dehydroabietyl)ethylenediamine), N-methyl-D-glucamines, N-methyl-D-glucamides, t-butyl amines, piperazine, phenylcyclohexyl-amine, choline, tromethamine, and salts with amino acids such as arginine, lysine and the like. Basic nitrogen-containing groups may be converted to an ammonium ion or quaternized with agents such as lower alkyl halides (e.g. methyl, ethyl, propyl, and butyl chlorides, bromides and iodides), dialkyl sulfates (e.g. dimethyl, diethyl, dibutyl, and diamyl sulfates), long chain halides (e.g. decyl, lauryl, myristyl and stearyl chlorides, bromides and iodides), aralkyl halides (e.g. benzyl and phenethyl bromides), and others.

The term "pharmaceutically acceptable anion" refers to an anion suitable for forming a pharmaceutically acceptable salt.

Further examples of pharmaceutically acceptable salts that may be used with the instant invention include, but are not limited to, fluoride, chloride, bromide and iodide.

In general, salts of compounds are intended to be pharmaceutically acceptable salts within the scope of the invention.

The term "purified", "in purified form" or "in isolated and purified form" for a compound refers to the physical state of said compound after being isolated from a synthetic process or natural source or combination thereof. Thus, the term "purified", "in purified form" or "in isolated and purified form" for a compound refers to the physical state of

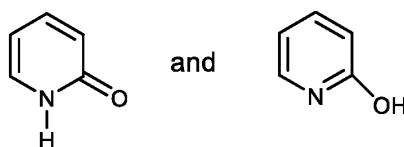
said compound after being obtained from a purification process or processes described herein or well known to the skilled artisan, and in sufficient purity to be characterized by standard analytical techniques described herein or well known to the skilled artisan.

Compounds of the invention include any form of the compound including *in situ* in a

5 reaction mixture as well as in isolated and purified form obtained by routine techniques.

Also included are polymorphic forms of the compounds of the invention and solvates and prodrugs thereof.

Certain compounds of the invention may exist in different tautomeric forms, for example, but are not limited to, ketone/enol tautomeric forms, imine-enamine tautomeric
10 forms, and for example heteroaromatic forms such as the following moieties:



In the same manner, unless indicated otherwise, presenting a structural representation of any tautomeric form of a compound which exhibits tautomerism is meant to include all such tautomeric forms of the compound. Accordingly, where compounds of the invention, their
15 salts, and solvates and prodrugs thereof, may exist in different tautomeric forms or in equilibrium among such forms, all such forms of the compound are embraced by, and included within the scope of the invention.

In another aspect, the present invention provides pharmaceutical compositions comprising one or more compounds of the invention. As used herein, the term

20 **"pharmaceutical composition" comprises at least one pharmaceutically active compound** and at least one excipient, and is intended to encompass both the combination of the specified ingredients in the specified amounts, and any product which results, directly or indirectly, from combination of the specified ingredients in the specified amounts.

As will be appreciated by the ordinarily skilled artisan, excipients are any constituent which adapts the composition to a particular route of administration or aids the processing of a composition into a dosage form without itself exerting an active pharmaceutical effect. In general compositions comprise more than one excipient depending upon the route of administration and the characteristics of the active being administered. Examples of excipients which impart to the composition properties which make it easier to handle or process include, but are not limited to, lubricants or pressing aids in powdered medicaments intended to be tableted, and emulsion stabilizers in compositions in which the active is present in the form of an emulsion. Examples of excipients which adapt a composition to a desired route of administration are, for example, but not limited to, for oral administration, absorption enhancers promoting absorption from the gastrointestinal tract, for transdermal or transmucosal administration, penetration enhancers, for example, those employed in adhesive skin "patch" or compositions for buccal administration.

Notwithstanding the function excipients perform in a composition, excipients are collectively termed herein "a carrier." Typically, formulations may comprise up to about 95 percent active ingredient and the balance carrier, although formulations with different ratios may be prepared. In general, acceptable pharmaceutical compositions contain a suitable concentration of the active that an effective amount of the PCSK9 antagonist can be provided in an individual dosage form of acceptable volume based upon the route of administration such that it can provide a therapeutic serum level of the active for an acceptable period of time in a subject to whom the composition is administered and the composition will retain biological activity during storage within an acceptable temperature range for an acceptable period of time.

Pharmaceutical composition, as used herein, refers both to a bulk composition, that is, formulated material that has not yet been formed into individual dosage units for administration, and the composition contained within individual dosage units.

While compositions of the invention may be employed in bulk form, it will be appreciated that for most applications compositions will be incorporated into a dosage form providing individual units suitable for administration to a patient, each dosage form comprising an amount of the selected composition which contains an effective amount of said one or more compounds of Formula I. Examples of suitable dosage forms include, but are not limited to, dosage forms adapted for: (i) oral administration, e.g., a liquid, gel, powder, solid or semi-solid pharmaceutical composition which is loaded into a capsule or pressed into a tablet and may comprise additionally one or more coatings which modify its release properties, for example, coatings which impart delayed release or formulations which have extended release properties; (ii) a dosage form adapted for administration through tissues of the oral cavity, for example, a rapidly dissolving tablet, a lozenge, a solution, a gel, a sachet or a needle array suitable for providing intramucosal administration; (iii) a dosage form adapted for administration via the mucosa of the nasal or upper respiratory cavity, for example a solution, suspension or emulsion formulation for dispersion in the nose or airway; (iv) a dosage form adapted for transdermal administration, for example, a patch, cream or gel; (v) a dosage form adapted for intradermal administration, for example, a microneedle array; (vi) a dosage form adapted for intravenous (IV) infusion, for example, over a prolonged period using an I.V. infusion pump; (vii) a dosage form adapted for intramuscular administration (IM), for example, an injectable solution or suspension, and which may be adapted to form a depot having extended release properties; (viii) a dosage form adapted for drip intravenous administration (IV), for example, a solution or suspension, for example, as an IV solution or a concentrate to be

injected into a saline IV bag; (ix) a dosage form adapted for subcutaneous administration, including administration over an extended time period by implanting a rod or other device which diffuses the compound into the surround tissue and thereby provides a continuous serum therapeutic level; or (x) a dosage form adapted for delivery via rectal or vaginal
5 mucosa, for example, a suppository.

Pharmaceutical compositions can be solid, semi-solid or liquid. Solid, semi-solid and liquid form preparations can be adapted to a variety of modes of administration, examples of which include, but are not limited to, powders, dispersible granules, mini-tablets, beads, which can be used, for example, for tableting, encapsulation, or direct administration. In
10 addition, liquid form preparations include, but are not limited to, solutions, suspensions and emulsions which for example, but not exclusively, can be employed in the preparation of formulations intended for ingestion, inhalation or intravenous administration (IV), for example, but not limited to, administration via drip IV or infusion pump, intramuscular injection (IM), for example, of a bolus which is released over an extended duration, direct IV
15 injection, or adapted to subcutaneous routes of administration.

Other routes of administration which may be contemplated include intranasal administration, or for administration to some other mucosal membrane. Formulations prepared for administration to various mucosal membranes may also include additional components adapting them for such administration, for example, viscosity modifiers.

20 Although in some embodiments, compositions suitable for use in a solid oral dosage form, for example, a tablet or quick-melt mouth-dissolving formulation are preferable routes of administration for a compound of the invention or a salt thereof, a composition of the invention may be formulated for administration via other routes mentioned above.

Examples include aerosol preparations, for example, suitable for administration via
25 inhalation or via nasal mucosa, may include solutions and solids in powder form, which may

be in combination with a pharmaceutically acceptable propellant, for example, an inert compressed gas, e.g. nitrogen. Also included are solid form preparations which are intended to be converted, shortly before use, to a suspension or a solution, for example, for oral or parenteral administration. Examples of such solid forms include, but are not limited
5 to, freeze dried formulations and liquid formulations adsorbed into a solid absorbent medium.

For example, the compounds of the invention may also be deliverable transdermally or transmucosally, for example, from a liquid, suppository, cream, foam, gel, or rapidly dissolving solid form. It will be appreciated that transdermal compositions can take also the
10 form of creams, lotions, aerosols and/or emulsions and can be provided in a unit dosage form which includes a transdermal patch of any known in the art, for example, a patch which incorporates either a matrix comprising the pharmaceutically active compound or a reservoir which comprises a solid or liquid form of the pharmaceutically active compound.

Examples of pharmaceutically acceptable carriers and methods of manufacture for
15 various compositions mentioned above may be found in A. Gennaro (ed.), Remington: The Science and Practice of Pharmacy, 20th Edition, (2000), Lippincott Williams & Wilkins, Baltimore, MD. Additional examples of publications addressing formulation issues may be found in: Pharmaceutical compositions may be formulated by any number of strategies known in the art, see, e.g., McGoff and Scher, 2000 *Solution Formulation of*
20 *Proteins/Peptides*: In – McNally, E.J., ed. Protein Formulation and Delivery. New York, NY: Marcel Dekker; pp. 139-158; Akers & Defilippis, 2000, *Peptides and Proteins as Parenteral Solutions*. In – Pharmaceutical Formulation Development of Peptides and Proteins. Philadelphia, PA: Taylor and Francis; pp. 145-177; Akers *et al.*, 2002, *Pharm. Biotechnol.* 14:47-127.

In another aspect the present invention provides methods of employing PCSK9-specific antagonist compounds described herein for antagonizing PCSK9 function; said **methods of which are further described below. Use of the term "antagonizing" throughout** the present application refers to providing to the affected tissue(s) a substance which

5 opposes the action of, inhibits, counteracts, neutralizes or curtails one or more functions of PCSK9 in the affected tissues. Inhibition or antagonism of one or more of PCSK9-associated functional properties can be readily determined according to methodologies known to the art (see, e.g., Barak & Webb, 1981 *J. Cell Biol.* 90:595-604; Stephan & Yurachek, 1993 *J. Lipid Res.* 34:325330; and McNamara *et al.*, 2006 *Clinica Chimica Acta*

10 369:158-167) as well as those described herein. Inhibition or antagonism will effectuate a decrease in PCSK9 activity relative to that seen in the absence of the antagonist or, for example, that seen relative to the activity observed when a control antagonist of irrelevant specificity is present. Preferably, a PCSK9-specific antagonist in accordance with the present invention antagonizes PCSK9 functioning to the point that there is a decrease of at

15 least 10%, of the measured parameter including but not limited to the activities disclosed herein, and more preferably, a decrease of at least 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90% and 95% of the measured parameter. Such inhibition/antagonism of PCSK9 functioning is particularly effective in those instances where PCSK9 functioning is contributing at least in part to a particular phenotype, disease, disorder or condition which is

20 negatively impacting the subject.

In one aspect, the present invention provides a method for antagonizing the activity of PCSK9, which comprises contacting a cell, population of cells or tissue sample capable of being affected by PCSK9 (*i.e.*, which expresses and/or comprises LDL receptors) with a PCSK9-specific antagonist disclosed herein under conditions that allow said antagonist to

25 bind to PCSK9 when present and inhibit PCSK9's inhibition of cellular LDL uptake. In some

embodiments of the present invention include such methods wherein the cell is a human cell. Additional embodiments of the present invention include such methods wherein the cell is a murine cell.

In one aspect, the present invention provides a method for antagonizing the activity of PCSK9 in a subject, which comprises administering to the subject a therapeutically effective amount of a PCSK9-specific antagonist of the present invention. In some embodiments, the methods for antagonizing PCSK9 function are for the treatment, as defined herein, of a PCSK9-associated disease, disorder or condition or, alternatively, for providing therapy in a disease, disorder or condition that could benefit from the effects of a PCSK9 antagonist.

The present invention, thus, contemplates the use of PCSK9-specific antagonists described herein in various methods of treatment where antagonizing PCSK9 function is desirable. As used herein, the term "method of treatment" **relates to a course of action** resulting in a change in at least one symptom of a disease state which can be prophylactic or therapeutic in nature. In some embodiments, the present invention relates to a method of treatment for a condition associated with and/or attributed to PCSK9 activity, or a condition where the functioning of PCSK9 is contraindicated for a particular subject, the method comprising administering to the subject a therapeutically effective amount of a PCSK9- antagonist compound of Formula I, or pharmaceutically acceptable salt thereof. In some embodiments, the condition may be atherosclerosis, hypercholesterolemia, coronary heart disease, metabolic syndrome, acute coronary syndrome or related cardiovascular disease and cardiometabolic conditions, or may be a disease state or condition in which PCSK9 activity is contraindicated.

Methods of treatment in accordance with the present invention comprise administering to an individual a therapeutically (or prophylactically) effective amount of a

PCSK9-specific antagonist of the present invention. Use of the terms "therapeutically effective" or "prophylactically effective" in reference to an amount refers to the amount necessary at the intended dosage to achieve the desired therapeutic and/or prophylactic effect for the period of time desired. The desired effect may be, for example, the alleviation, amelioration, reduction or cessation of at least one symptom associated with the treated condition. These amounts will vary, as the skilled artisan will appreciate, according to various factors, including but not limited to the disease state, age, sex, and weight of the individual, and the ability of the PCSK9-specific antagonist to elicit the desired effect in the individual. The response may be documented by *in vitro* assay, *in vivo* non-human animal studies, and/or further supported from clinical trials.

In some embodiments it is preferred to administer a PCSK9 antagonist compound of the invention in the form of a pharmaceutical composition as described herein.

Dosing of antagonist therapeutics is well within the realm of the skilled artisan, see, e.g., Lederman *et al.*, 1991 *Int. J. Cancer* 47:659-664; Bagshawe *et al.*, 1991 *Antibody, Immunoconjugates and Radiopharmaceuticals* 4:915-922, and will vary based on a number of factors, for example, but not limited to, those mentioned above, including the condition of the patient, the area being treated, the route of administration, and the treatment desired, for example, prophylaxis or acute treatment and the like. A physician or veterinarian of ordinary skill can readily determine and prescribe the effective therapeutic amount of the antagonist.

The subject may be in need of, or desire, treatment for an existing disease or medical condition. As used herein, the subject "in need" of treatment of an existing condition encompasses both a determination of need by a medical professional as well as the desire of the subject for such treatment. When a compound or a salt thereof is provided in combination with one or more other active agents, "administration" and its variants are

each understood to include provision of the compound or its salt and the other agents contemporaneously or simultaneously or over a course of separate administrations over a period of time. When the agents of a combination are administered at the same time, they can be administered together in a single composition or they can be administered

5 separately. It is understood that a "combination" of active agents can be a single composition containing all of the active agents or multiple compositions each containing one or more of the active agents. In the case of two active agents a combination can be either a single composition comprising both agents or two separate compositions each comprising one of the agents; in the case of three active agents a combination can be
10 either a single composition comprising all three agents, three separate compositions each comprising one of the agents, or two compositions one of which comprises two of the agents and the other comprises the third agent; and so forth.

The compositions and combinations of the present invention are suitably administered in effective amounts. The term "effective amount" means the amount of active
15 compound sufficient to antagonize PCSK9 and thereby elicit the response being sought (*i.e.*, induce a therapeutic response in the treatment or management of conditions associated with or impacted by PCSK9 function, including, but not limited to atherosclerosis, hypercholesterolemia, coronary heart disease, metabolic syndrome, acute coronary syndrome, and related cardiovascular disease and cardiometabolic conditions in
20 an animal or human).

The actual dosage employed may be varied depending upon the requirements of the patient and the severity of the condition being treated. Determination of the proper dosage regimen for a particular situation is within the skill in the art, for example, as described in the standard literature, for example, as described in the "Physicians' Desk Reference" (PDR),
25 e.g., 1996 edition (Medical Economics Company, Montvale, NJ 07645-1742, USA), the

Physician's Desk Reference, 56th Edition, 2002 (published by Medical Economics company, Inc. Montvale, NJ 07645-1742), or **the Physician's Desk Reference**, 57th Edition, 2003 (published by Thompson PDR, Montvale, NJ 07645-1742); the disclosures of which is incorporated herein by reference thereto. For convenience, the total daily dosage may be
5 divided and administered in portions during the day as required or delivered continuously.

The PCSK9-specific antagonist may be administered to an individual by any route of administration appreciated in the art, including but not limited to oral administration, administration by injection (specific embodiments of which include intravenous, subcutaneous, intraperitoneal or intramuscular injection), or administration by inhalation,
10 intranasal, or topical administration, either alone or in combination with other agents designed to assist in the treatment of the individual. The PCSK9-specific antagonist may also be administered by injection devices, injector pens, needleless devices; and subcutaneous patch delivery systems. The route of administration should be determined based on a number of considerations appreciated by the skilled artisan including, but not
15 limited to, the desired physiochemical characteristics of the treatment.

One or more additional pharmacologically active agents may be administered in combination with a compound of Formula I. An additional active agent (or agents) is intended to mean a pharmaceutically active agent (or agents) that is active in the body, including pro-drugs that convert to pharmaceutically active form after administration, which
20 are different from the compound of Formula I, and also includes free-acid, free-base and pharmaceutically acceptable salts of said additional active agents. Generally, any suitable additional active agent or agents, including but not limited to anti-hypertensive agents, anti-atherosclerotic agents such as a lipid modifying compound, anti-diabetic agents and/or anti-obesity agents may be used in any combination with the compound of Formula I in a single
25 dosage formulation (a fixed dose drug combination), or may be administered to the subject

in one or more separate dosage formulations which allows for concurrent or sequential administration of the active agents (co-administration of the separate active agents).

Examples of additional active agents which may be employed include but are not limited to angiotensin converting enzyme inhibitors (e.g., alacepril, benazepril, captopril, 5 ceronapril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, imidapril, lisinopril, moveltipril, perindopril, quinapril, ramipril, spirapril, temocapril, ortrandolapril), angiotensin II receptor antagonists (e.g., losartan i.e., COZAAR®, valsartan, candesartan, olmesartan, telmesartan and any of these drugs used in combination with hydrochlorothiazide such as HYZAAR®); neutral endopeptidase inhibitors (e.g., thiorphan and phosphoramidon), aldosterone 10 antagonists, aldosterone synthase inhibitors, renin inhibitors (e.g. urea derivatives of di- and tri-peptides (See U.S. Pat. No. 5,116,835), amino acids and derivatives (U.S. Patents 5,095,119 and 5,104,869), amino acid chains linked by non-peptidic bonds (U.S. Patent 5,114,937), di- and tri-peptide derivatives, peptidyl amino diols and peptidyl beta-aminoacyl aminodiols carbamates, and small molecule renin inhibitors (including diol sulfonamides and 15 sulfinyls), N-morpholino derivatives, N-heterocyclic alcohols and pyroimidazolones; also, pepstatin derivatives and fluoro- and chloro-derivatives of statone-containing peptides, enalkrein, RO 42-5892, A 65317, CP 80794, ES 1005, ES 8891, SQ 34017, aliskiren (2(S),4(S),5(S),7(S)-N-(2-carbamoyl-2-methylpropyl)-5-amino-4-hydroxy-2,7-diisopropyl-8-[4-methoxy-3-(3-methoxypropoxy)-phenyl]-octanamid hemifumarate) SPP600, SPP630 and 20 SPP635), endothelin receptor antagonists, phosphodiesterase-5 inhibitors (e.g. sildenafil, tadalafil and vardenafil), vasodilators, calcium channel blockers (e.g., amlodipine, nifedipine, verapamil, diltiazem, gallopamil, niludipine, nimodipins, nicardipine), potassium channel activators (e.g., nicorandil, pinacidil, cromakalim, minoxidil, aprilkalim, loprazolam), diuretics (e.g., hydrochlorothiazide), sympatholitics, beta-adrenergic blocking drugs (e.g., 25 propranolol, atenolol, bisoprolol, carvedilol, metoprolol, or metoprolol tartate), alpha

adrenergic blocking drugs (e.g., doxazocin, prazosin or alpha methyldopa) central alpha adrenergic agonists, peripheral vasodilators (e.g. hydralazine); lipid lowering agents e.g., HMG-CoA reductase inhibitors such as simvastatin and lovastatin which are marketed as ZOCOR® and MEVACOR® in lactone pro-drug form and function as inhibitors after

5 administration, and pharmaceutically acceptable salts of dihydroxy open ring acid HMG-CoA reductase inhibitors such as atorvastatin (particularly the calcium salt sold in LIPITOR®), rosuvastatin (particularly the calcium salt sold in CRESTOR®), pravastatin (particularly the sodium salt sold in PRAVACHOL®), fluvastatin (particularly the sodium salt sold in LESCOL®), crivastatin, and pitavastatin; a cholesterol absorption inhibitor such as

10 ezetimibe (ZETIA®) and ezetimibe in combination with any other lipid lowering agents such as the HMG-CoA reductase inhibitors noted above and particularly with simvastatin (VYTORIN®) or with atorvastatin calcium; niacin in immediate-release or controlled release forms and/or with an HMG-CoA reductase inhibitor; niacin receptor agonists such as acipimox and acifran, as well as niacin receptor partial agonists; metabolic altering agents

15 including insulin and insulin mimetics (e.g., insulin degludec, insulin glargine, insulin lispro), dipeptidyl peptidase-IV (DPP-4) inhibitors (e.g., sitagliptin, alogliptin, omarigliptin, linagliptin, vildagliptin); **insulin sensitizers, including (i) PPAR γ agonists, such as the glitazones (e.g. pioglitazone, AMG 131, MBX2044, mitoglitazone, lobeglitazone, IDR-105, rosiglitazone, and balaglitazone), and other PPAR ligands, including (1) PPAR α/γ dual agonists (e.g.,**

20 **ZYH2, ZYH1, GFT505, chiglitazar, muraglitazar, aleglitazar, sodelglitazar, and naveglitazar); (2) PPAR α agonists such as fenofibric acid derivatives (e.g., gemfibrozil, clofibrate, ciprofibrate, fenofibrate, bezafibrate), (3) selective PPAR γ modulators (SPPAR γ M's), (e.g., such as those disclosed in WO 02/060388, WO 02/08188, WO 2004/019869, WO 2004/020409, WO 2004/020408, and WO 2004/066963); and (4) PPAR γ**

25 partial agonists; (ii) biguanides, such as metformin and its pharmaceutically acceptable

salts, in particular, metformin hydrochloride, and extended-release formulations thereof, such as **Glumetza™, Fortamet™, and GlucophageXR™**; and (iii) protein tyrosine phosphatase-1B (PTP-1B) inhibitors (e.g., ISIS-113715 and TTP814); insulin or insulin analogs (e.g., insulin detemir, insulin glulisine, insulin degludec, insulin glargine, insulin lispro and inhalable formulations of each); leptin and leptin derivatives and agonists; amylin and amylin analogs (e.g., pramlintide); sulfonylurea and non-sulfonylurea insulin secretagogues (e.g., tolbutamide, glyburide, glipizide, glimepiride, mitiglinide, meglitinides, nateglinide and repaglinide); α -glucosidase inhibitors (e.g., acarbose, voglibose and miglitol); glucagon receptor antagonists (e.g., MK-3577, MK-0893, LY-2409021 and KT6-971); incretin mimetics, such as GLP-1, GLP-1 analogs, derivatives, and mimetics; and GLP-1 receptor agonists (e.g., dulaglutide, semaglutide, albiglutide, exenatide, liraglutide, lixisenatide, taspoglutide, CJC-1131, and BIM-51077, including intranasal, transdermal, and once-weekly formulations thereof); bile acid sequestering agents (e.g., colestilan, colestimide, colesevalam hydrochloride, colestipol, cholestyramine, and dialkylaminoalkyl derivatives of a cross-linked dextran), acyl CoA:cholesterol acyltransferase inhibitors, (e.g., avasimibe); antiobesity compounds; agents intended for use in inflammatory conditions, such as aspirin, non-steroidal anti-inflammatory drugs or NSAIDs, glucocorticoids, and selective cyclooxygenase-2 or COX-2 inhibitors; glucokinase activators (GKAs) (e.g., **AZD6370**); **inhibitors of 11 β -hydroxysteroid dehydrogenase type 1**, (e.g., such as those disclosed in U.S. Patent No. 6,730,690, and LY-2523199); CETP inhibitors (e.g., anacetrapib, torcetrapib, and evacetrapib); inhibitors of fructose 1,6-bisphosphatase, (e.g., such as those disclosed in U.S. Patent Nos. 6,054,587; 6,110,903; 6,284,748; 6,399,782; and 6,489,476); inhibitors of acetyl CoA carboxylase-1 or 2 (ACC1 or ACC2); AMP-activated Protein Kinase (AMPK) activators; other agonists of the G-protein-coupled receptors: (i) GPR-109, (ii) GPR-119 (e.g., MBX2982 and PSN821), and (iii) GPR-40 (e.g.,

TAK875); SSTR3 antagonists (e.g., such as those disclosed in WO 2009/001836); neuromedin U receptor agonists (e.g., such as those disclosed in WO 2009/042053, including, but not limited to, neuromedin S (NMS)); SCD modulators; GPR-105 antagonists (e.g., such as those disclosed in WO 2009/000087); SGLT inhibitors (e.g., ASP1941, 5 SGLT-3, empagliflozin, dapagliflozin, canagliflozin, BI-10773, ertugliflozin, remogliflozin, TS-071, tofogliflozin, ipragliflozin, and LX-4211); inhibitors of acyl coenzyme A:diacylglycerol acyltransferase 1 and 2 (DGAT-1 and DGAT-2); inhibitors of fatty acid synthase; inhibitors of acyl coenzyme A:monoacylglycerol acyltransferase 1 and 2 (MGAT-1 and MGAT-2); agonists of the TGR5 receptor (also known as GPBAR1, BG37, GPCR19, 10 GPR131, and M-BAR); ileal bile acid transporter inhibitors; PACAP, PACAP mimetics, and PACAP receptor 3 agonists; PPAR agonists; protein tyrosine phosphatase-1B (PTP-1B) inhibitors; IL-1b antibodies, (e.g., XOMA052 and canakinumab); and bromocriptine mesylate and rapid-release formulations thereof; or with other drugs beneficial for the treatment of the above-mentioned conditions or disorders including the free-acid, free-base, 15 and pharmaceutically acceptable salt forms of the above active agents where chemically possible.

The compounds of the present invention can be readily prepared according to the following reaction schemes and examples, or modifications thereof, using readily available starting materials, reagents and conventional synthesis procedures. In these reactions, it is 20 also possible to make use of known variants. For purification of the compounds using reverse phase chromatography (either HPLC or MPLC, as noted below), a C18 column was used. Other methods for preparing compounds of the invention will be readily apparent to the person of ordinary skill in the art in light of the following reaction schemes and examples. Abbreviations listed below may be used in the exemplary schemes and/or 25 examples herein.

- ACN is acetonitrile
- AcOH is acetic acid
- AcO·NH₄ is ammonium acetate
- Boc₂O is di-*tert*-butyl dicarbonate
- 5 Bn is benzyl
- BnBr is benzyl bromide
- BzCl is benzoyl chloride
- CBr₄ is tetrabromomethane
- Cbz-Cl is benzyl chloroformate
- 10 DBU is 1,8-Diazabicyclo[5.4.0]undec-7-ene
- DCC is dicyclohexylcarbodiimide
- DCE is 1,2-dichloroethane
- DCM is dichloromethane
- DEA is N,N-diethylamine
- 15 DIAD is (E)-diisopropyl diazene-1,2-dicarboxylate
- DIEA or DIPEA is N,N-diisopropylethylamine
- DMAP is 4-dimethylaminopyridine
- DMF is N,N-dimethylformamide
- DMSO is dimethyl sulfoxide
- 20 EA or EtOAc is ethyl acetate
- EtOH is ethanol
- Et₂O is diethyl ether
- Fmoc is fluorenylmethyloxycarbonyl protecting group
- Fmoc-Cl is (9*H*-fluoren-9-yl)methyl carbonochloridate
- 25 Fmoc-D-Dap(Boc)-OH is N-α-(9-Fluorenylmethyloxycarbonyl)-N-β-*t*-butyloxycarbonyl-D-2,3-diaminopropionic acid

Fmoc-Osu is Fmoc N-hydroxysuccinimide ester

HATU is 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate

HPLC is High Performance Liquid Chromatography

5 IPA is isopropyl alcohol

LiOH is lithium hydroxide

LC/MS is Liquid chromatography--mass spectrometry

Me₃N is trimethyl amine

MeOH is methanol

10 MPLC is Medium pressure liquid chromatography

MsCl is methanesulfonyl chloride

NaBH(OAc)₃ is sodium triacetoxyborohydride

NMR is Nuclear Magnetic Resonance

NsCl is 4-nitrobenzene-1-sulfonyl chloride

15 PE is petroleum ether

Pd₂(dba)₃(HCCl₃) is tris(dibenzylideneacetone)dipalladium(0)-chloroform adduct

PPh₃ is triphenylphosphine

PdCl₂(dppf) or Pd(ii)(dppf)Cl₂ is dichloro[1,1'-bis(diphenylphosphino)ferrocene]palladium(II)

Pd(dppf)Cl₂CH₂Cl₂ is dichloro[1,1'-bis(diphenylphosphino)ferrocene]palladium(II)

20 dichloromethane adduct

Pd(PPh₃)₄ is tetrakis(triphenylphosphine)palladium

PPT_s is pyridinium p-toluenesulfonate

[Rh(OAc)₂]₂ is rhodium(II) acetate dimer

RT or r.t. or rt is room temperature

25 tBuOAc is tert-butyl acetate

TEA is triethylamine

TFA is trifluoroacetic acid

TFE is tetrafluoroethylene

THF is tetrahydrofuran

Tf₂O is trifluoromethanesulfonic anhydride

Teoc-OSu is 2,5-dioxopyrrolidin-1-yl (2-(trimethylsilyl)ethyl) carbonate

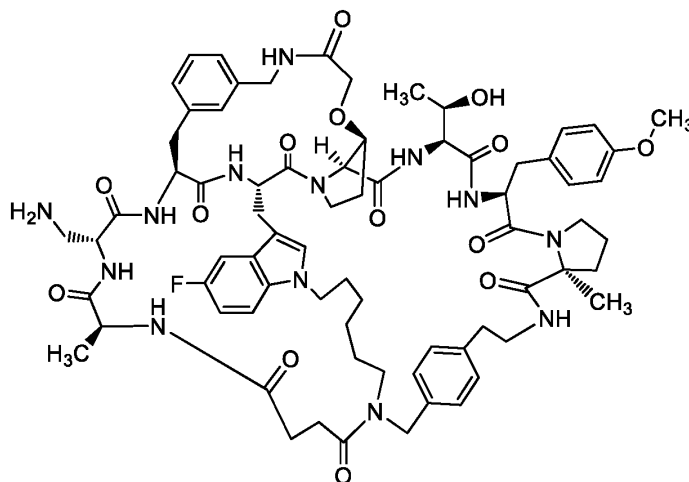
5 TBAF is tetrabutylammonium fluoride

TMS is tetramethylsilane

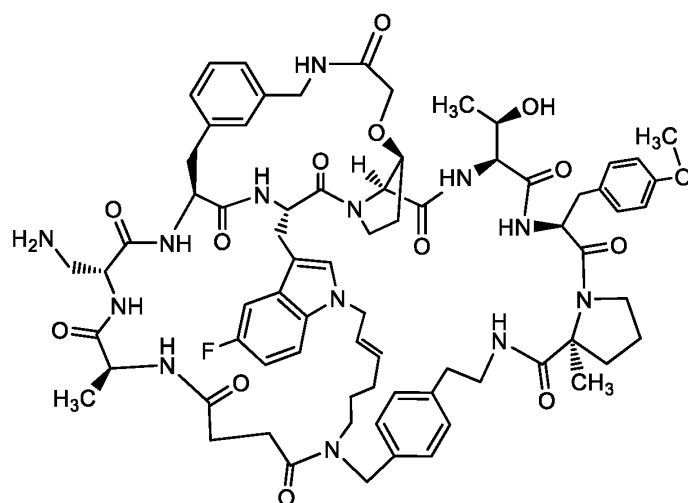
Zhan's catalyst 1B is dichloro(1,3-bis(2,4,6-trimethylphenyl)-2-imidazolidinyldene)((5-
((dimethylamino)sulfonyl)-2-(1-methylethoxy-O)phenyl)methylene-C)ruthenium(II)

[also described as 1,3-Bis(2,4,6-trimethylphenyl)-4,5-dihydroimidazol-2-ylidene[2-(i-
10 propoxy)-5-(N,N-dimethylaminosulfonyl)phenyl[methylenruthenium (II) dichloride]

EXAMPLE 1 Preparation of Ex-01 and Ex-51

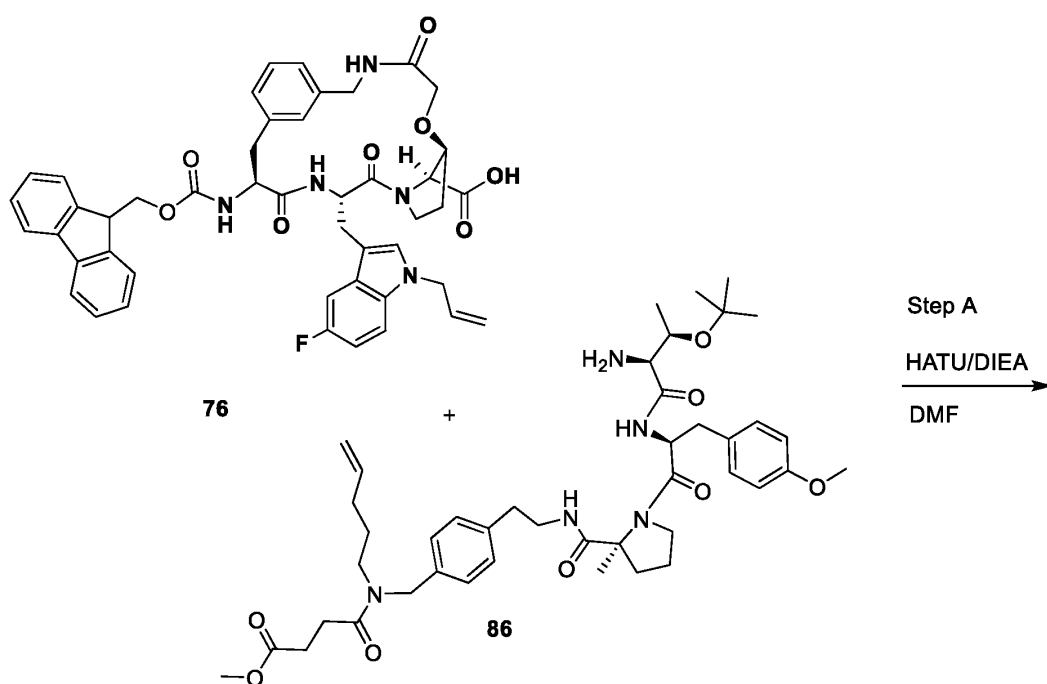


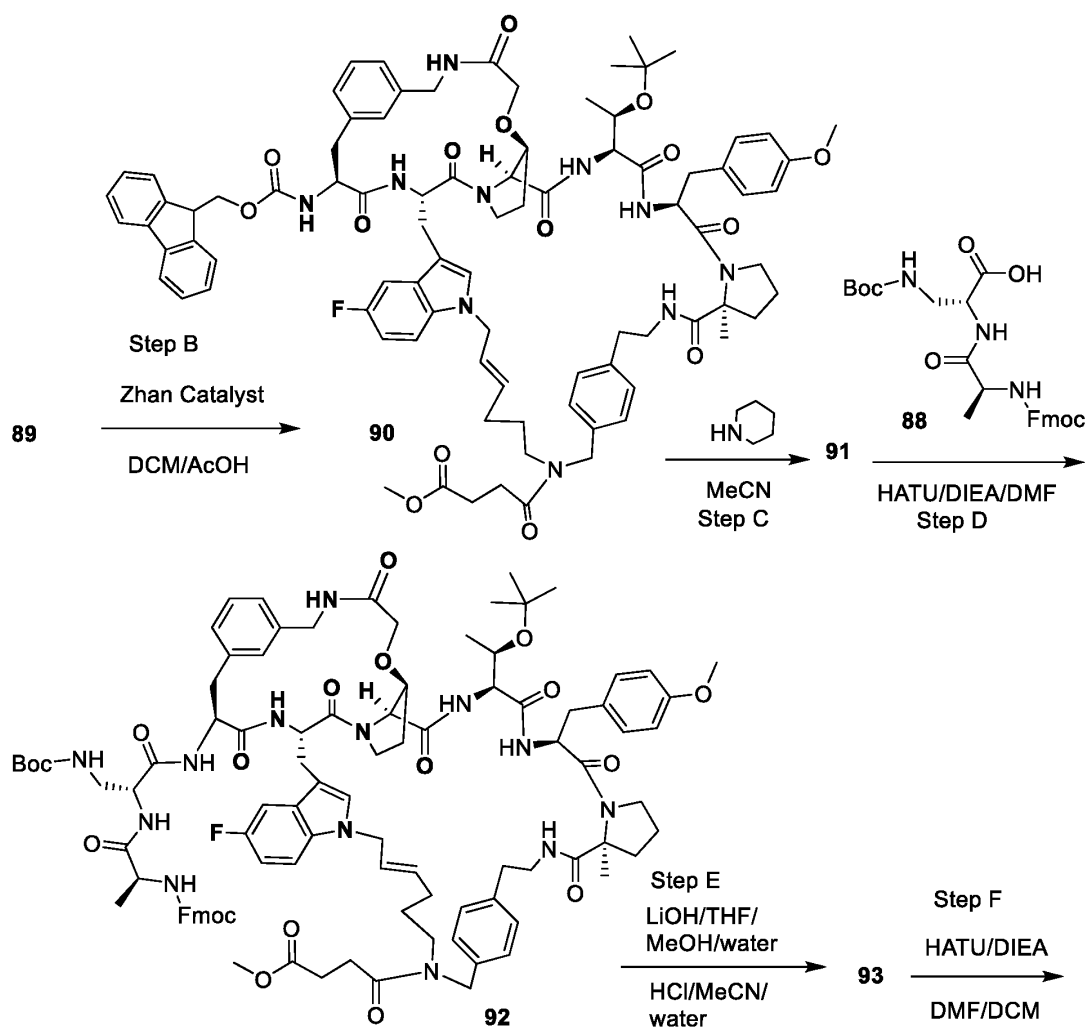
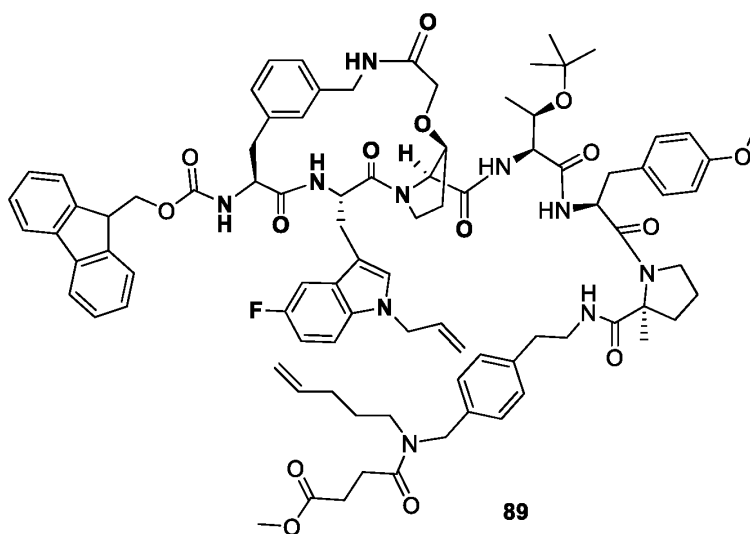
Ex-01, (free base illustrated)

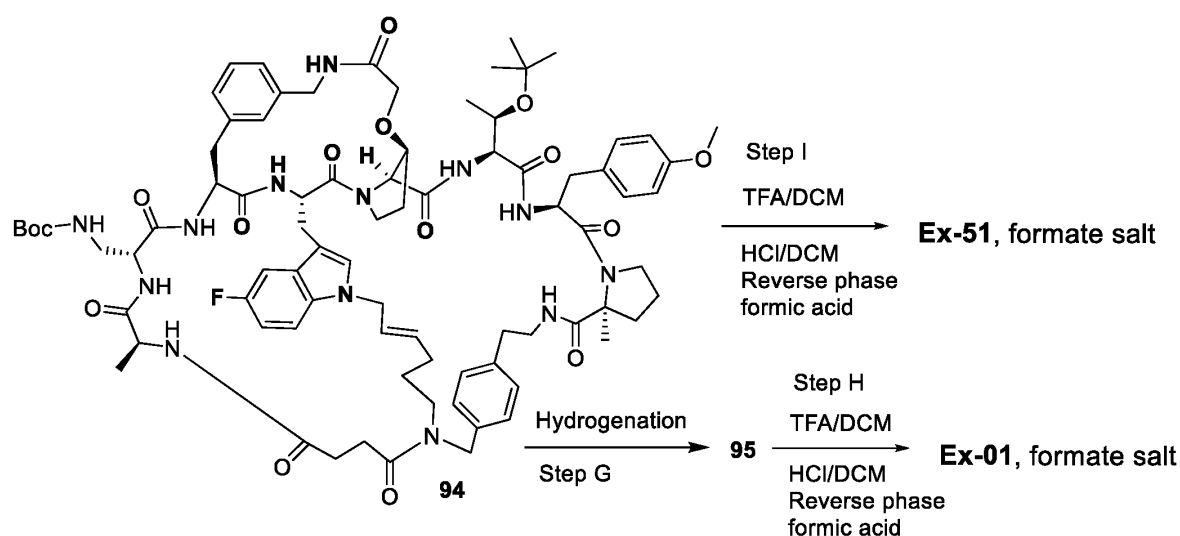


Ex-51, (free base illustrated)

Salt forms of compounds **Ex-01** and **Ex-51** were prepared from intermediates **76** and **86** (preparation following) in accordance with the following scheme:







Step A: Preparation of intermediate 89

To a solution of **76** (1.56 g, 1.917 mmol, preparation following) and **86** (1.506 g, 1.936 mmol, preparation following) in DMF (40 ml) was added HATU (0.802 g, 2.108 mmol) and DIEA (0.670 ml, 3.83 mmol). The resulting solution was stirred at rt for 50 min, then partitioned between EtOAc (300 mL) and brine (100 mL). The organic phase was washed with brine (2x100 mL), dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using MeOH/DCM as eluting solvents to give **89**. LC/MS: (M+1)⁺: 1573.8.

Step B: Preparation of intermediate 90

A solution of **89** (0.68 g, 0.432 mmol) in DCM (500 ml) and acetic acid (40 mL) was bubbled with N₂ for 20 min followed by addition of Zhan catalyst-1b (0.222 g, 0.302 mmol). The resulting mixture was further bubbled with N₂ for 20 min, then heated at 55 °C for 5 h. After cooling to rt the mixture was filtered through celite, the filtrate was concentrated, and the residue was purified on silica gel column using MeOH/DCM as eluting solvents to give **90** (cis/trans mixture). LC/MS: (M+1)⁺: 1545.7.

Step C: Preparation of intermediate 91

To a solution of **90** (cis/trans mixture) (65 mg, 0.042 mmol) in acetonitrile (2 ml) was added piperidine (0.042 ml, 0.420 mmol). The resulting solution was stirred at rt for 1 h, then

concentrated and the residue was dissolved in acetonitrile (4 mL) and concentrated again. The residue was further dried under high vacuum for 30 min to give **91** (cis and trans mixture). LC/MS: (M+1)⁺: 1324.0.

Step D: Preparation of intermediate 92

- 5 To a solution of **91** (cis/trans mixture) (548 mg, 0.414 mmol) and **88** (227 mg, 0.455 mmol, preparation following) in DMF (10 ml) at 0 °C was added HATU (181 mg, 0.476 mmol) and DIEA (0.166 ml, 0.952 mmol). The resulting solution was stirred at 0 °C for 1 h, and the solution was purified by reverse phase MPLC over C18 column using acetonitrile (0.05%TFA)/water (0.05%TFA) as eluting solvents to give **92** (cis/trans mixture). LC/MS:
- 10 (M+1)⁺: 1803.5.

Step E: Preparation of intermediate 93

- To a solution of **92** (700 mg, 0.388 mmol) in THF (20 ml), MeOH (6 ml), and water (6 ml) at 0 °C was added 1N aqueous LiOH (3.11 ml, 3.11 mmol) dropwise, and the resulting solution was stirred at 0 °C for 23 h. The solution was neutralized by addition of 1N HCl to
- 15 pH 7-8, the volatile was evaporated, the aqueous phase was acidified to pH 5, and the mixture was purified by reverse phase MPLC over C18 column using acetonitrile (0.05%TFA)/water (0.05%TFA) as eluting solvents to give **93** as TFA salt. To a solution of **93** TFA salt (427 mg, 0.257 mmol) in water (70 mL) and acetonitrile (70 ml) at 0 °C was added 0.1 N HCl (13.5 ml, 1.350 mmol) dropwise, the resulting solution was stirred at 0 °C
- 20 for 5 min, then lyophilized to give **93** as HCl salt. LC/MS: (M+1)⁺: 1567.1.

Step F: Preparation of intermediate 94

- To a solution of **93** as HCl salt (200 mg, 0.125 mmol) in DMF (30 mL) was added HATU (56.9 mg, 0.150 mmol). The resulting solution was stirred at rt for 30 min, then diluted with DCM (400 mL) followed by addition of DIEA (0.065 mL, 0.374 mmol). The resulting solution
- 25 was stirred at rt for 1h, the volatile was evaporated on rotary evaporator, and the resulting

DMF solution was purified by reverse phase MPLC using acetonitrile (0.05%TFA)/water (0.05%TFA) as eluting solvents to give **94**. LC/MS: (M+1)⁺: 1549.2.

Step G: Preparation of intermediate 95

To a solution of **94** (14 mg, 9.04 μ mol) in MeOH (20 ml) was added 10% Pd/C (1.924 mg, 1.808 μ mol) and the resulting mixture was subjected to hydrogenation at rt via H₂ balloon for 1h. The mixture was filtered through celite and the filtrate was concentrated to give intermediate compound **95**. LC/MS: (M+1)⁺: 1550.9.

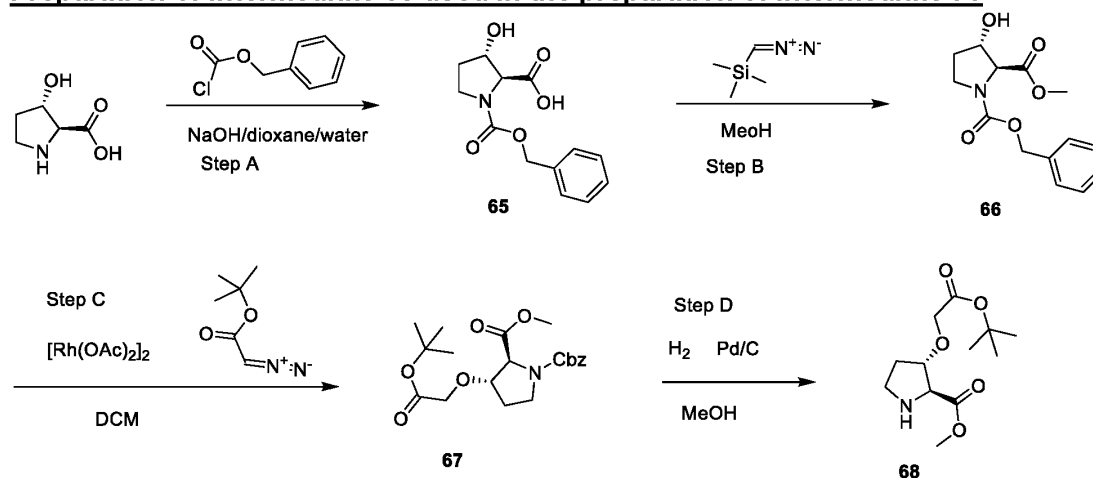
Step H: Preparation of Ex-01

Intermediate compound **95** prepared in the previous step (26 mg, 0.017 mmol) was dissolved in DCM (2 ml). To this solution was added TFA (6 mL, 78 mmol) and the resulting solution was stirred at rt for 30 min, then concentrated and the residue was dissolved in DCM (3 mL) and treated with 4N HCl in dioxane (0.042 mL, 0.168 mmol), and concentrated again to give **Ex-01** as a crude product. The crude **Ex-01** was purified by reverse phase HPLC using acetonitrile (0.1%formic acid)/water (0.1%formic acid) as eluting solvents to provide the formate salt form of **Ex-01**. LC/MS: (M+1)⁺: 1394.4.

Step I: Preparation of Ex-51

To a solution of intermediate compound **94** (30 mg, 0.019 mmol) in DCM (2 ml) was added TFA (4 ml, 51.9 mmol), and the reaction mixture was stirred at ambient temperature for 30 minutes, then concentrated. The residue was dissolved in DCM (2 mL), treated with HCl (4N in dioxane) (0.048 ml, 0.194 mmol), and concentrated to yield **Ex-51** as a HCl salt. The compound was purified by reverse phase HPLC using acetonitrile (0.1%formic acid)/water (0.1%formic acid) as mobile phase to provide the formate salt form of **Ex-51**. LC/MS: (M+1)⁺: 1392.0.

The following schemes and procedures were used to prepare intermediates **76**, **86** and **88** used in the procedures described above.

Preparation of Intermediate 68 used in the preparation of Intermediate 76**Step A: Preparation of Intermediate compound 65**

To a suspension of (2S,3S)-3-hydroxypyrrolidine-2-carboxylic acid (5.32 g, 40.6 mmol) in dioxane (100 ml) at 0 °C was added sodium hydroxide (122 ml, 122 mmol), followed by addition of benzyl chloroformate (6.50 ml, 44.6 mmol) dropwise. The resulting suspension was stirred at 0 °C for 5h. After removing the volatile, the aqueous phase was acidified to pH 3, then partitioned between 30%IPA/DCM(200 mL) and brine (50 mL), the aqueous phase was further extracted with 30%IPA/DCM(2x100mL). Combined organic phases were dried over Na₂SO₄ and concentrated to give (2S,3S)-1-((benzyloxy)carbonyl)-3-hydroxypyrrolidine-2-carboxylic acid (**65**). LC/MS: (M+1)⁺: 266.1.

Step B: Preparation of Intermediate compound 66

To a solution of **65** (7.48 g, 28.2 mmol) in MeOH (80 ml) was added TMS-Diazomethane (70.5 ml, 141 mmol) dropwise, and the resulting solution was stirred at rt for 10 min then quenched by addition of acetic acid (ca. 400 uL) dropwise. The solution was concentrated, and the residue was purified on silica gel column using EtOAc/hexane as eluting solvents to give **66**. LC/MS: (M+1)⁺: 280.1.

Step C: Preparation of Intermediate compound 67

A solution of **66** (4.81 g, 17.22 mmol) in DCM (200 mL) was bubbled with N₂ for 30 min, followed by addition of rhodium(ii) acetate dimer (0.761 g, 1.722 mmol). The mixture was cooled in a ice-water bath, and tert-butyl diazoacetate (3.58 mL, 25.8 mmol) were added at 0 °C dropwise. The resulting mixture was stirred at 0 °C for 1.5h. The reaction was

5 quenched by addition of water (100 mL), the mixture was extracted with DCM (3x100 mL), the combined organic phase was dried over Na₂SO₄, concentrated and the residue was purified by reverse phase MPLC using acetonitrile (0.05%TFA)/water (0.05%TFA) as eluting solvents. The fraction containing the product was concentrated and the aqueous

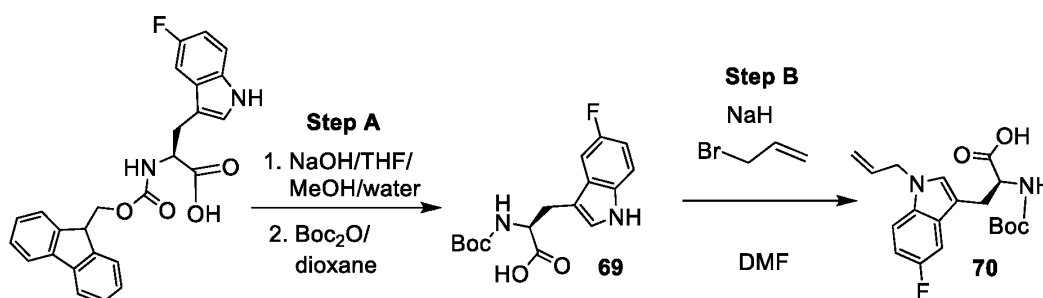
10 Na₂SO₄ and concentrated to give **67**. LC/MS: (M+1)⁺: 394.2.

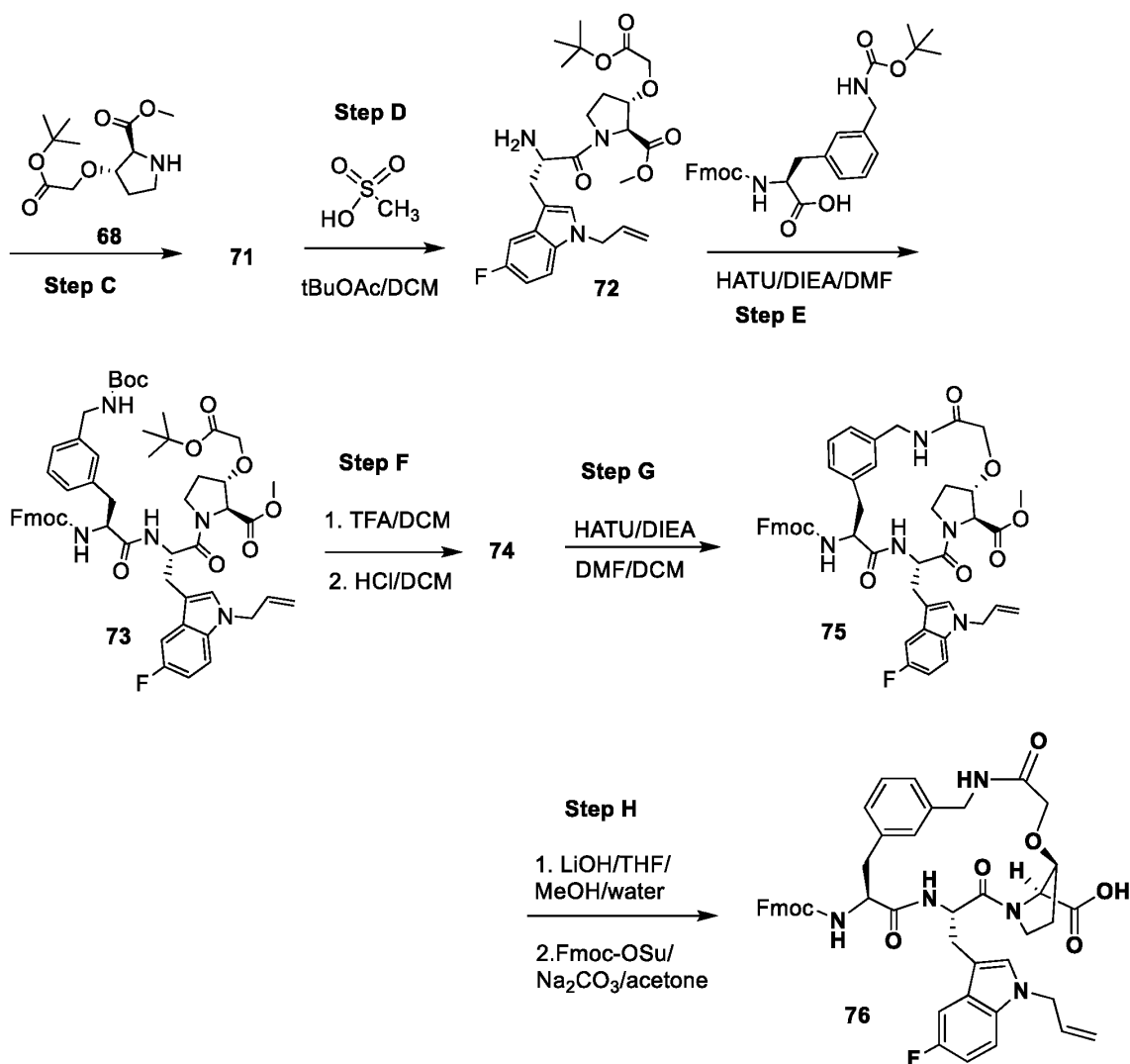
Step D: Preparation of Intermediate compound 68

To a solution of **67** (3.72 g, 9.46 mmol) in MeOH (80 ml) was added 10% Pd/C (0.805 g, 0.756 mmol) and the resulting mixture was subjected to hydrogenation via H₂ balloon at ambient temperature for 2 hours, then filtered through celite. The filtrate was concentrated

15 to give **68**. LC/MS: (M+1)⁺: 259.9.

Preparation of intermediate compound 76





Step A: Preparation of Intermediate 69

- 5 To a solution of (S)-2-((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(5-fluoro-1H-indol-3-yl)propanoic acid (3 g, 6.75 mmol) in THF (20 ml), MeOH (10 mL), and water (20.00 ml) at 0 °C was added NaOH (20.25 ml, 20.25 mmol) and the resulting solution was stirred at ambient temperature for 4 hours, then the volatile was evaporated. To the aqueous mixture was added dioxane (50 ml) and water (20 mL), the resulting solution was cooled to 0 °C
- 10 and Boc_2O (1.881 ml, 8.10 mmol) was added to the above solution. The resulting solution was stirred at 0 °C for 3 hours, the volatile was removed and the aqueous phase was

extracted with Et₂O (3x40 mL), acidified to pH 3, then extracted with DCM (3x100mL), followed by 30%IPA/DCM (2x80 mL). The combined organic phase was dried over Na₂SO₄ and concentrated to give **69**. LC/MS: (M+1)⁺: 322.9.

Step B: Preparation of Intermediate 70

- 5 To a solution of **69** (2.079 g, 6.45 mmol) in DMF (40 ml) at 0 °C was added 60% NaH in hexane (0.568 g, 14.19 mmol), and the resulting solution was stirred at 0 °C for 50 min followed by addition of allyl bromide (1.172 mL, 13.54 mmol) dropwise. The resulting solution was stirred at 0 °C for 1.5h, then quenched by addition of 1N HCl (ca. 3.68 mL). The solution was then partitioned between EtOAc (200 mL) and water (100 mL), the
10 organic phase was washed with brine (2x100 mL), dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using MeOH/DCM as eluting solvents to give **70**. LC/MS: (M+1)⁺: 363.0.

Step C: Preparation of Intermediate 71

- To a solution of **70** (2.239 g, 6.18 mmol) and **68** (1.842 g, 7.11 mmol) in DMF (30 ml) was
15 added HATU (2.82 g, 7.41 mmol) and DIEA (2.59 ml, 14.83 mmol), and the resulting solution was stirred at ambient temperature for 1 hour. The mixture was partitioned between EtOAc (200 mL) and brine (100 mL), the organic phase was washed with brine (3x100 mL), dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using EtOAc/hexane as eluting solvents to give **71**. LC/MS: (M+1)⁺: 604.2.

20 *Step D: Preparation of Intermediate 72*

To a solution of **71** (2.83 g, 4.69 mmol) in CH₂Cl₂ (20 ml) and tBuOAc (30 ml) at 0 °C was added methanesulfonic acid (1.218 ml, 18.75 mmol), and the resulting solution was stirred at 0 °C for 16.5 h, then ambient temperature for 2.5 h. The solution (**72**) was directly used in the next step. LC/MS: (M+1)⁺: 504.2.

25 *Step E: Preparation of Intermediate 73*

To a solution of (S)-2-((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(3-((((tert-butoxycarbonyl)amino)methyl)phenyl)propanoic acid (2.66 g, 5.16 mmol) in DMF (10 ml) was added HATU (1.961 g, 5.16 mmol) and DIEA (5.32 ml, 30.5 mmol), and the resulting solution was stirred at rt for 30 min then added to an ice-cold bath of the above prepared **72** solution. The resulting solution was stirred at ambient temperature for 1 hour. Volatiles were evaporated on rotary evaporator, and the residue was purified by reverse phase MPLC using acetonitrile (0.05%TFA)/water (0.05%TFA) as eluting solvents. Collected fractions were concentrated on rotary evaporator to give **73**. LC/MS: (M+1)⁺: 1002.1.

Step F: Preparation of Intermediate 74

To a solution of **73** (3.235 g, 3.23 mmol) in DCM (4 ml) was added TFA (7.46 ml, 97 mmol), and the resulting solution was stirred at ambient temperature for 1 hour, then concentrated. The residue was dissolved in DCM (10 mL), treated with 4N HCl in dioxane (3.23 ml, 12.91 mmol), then concentrated and the residue was dissolved in acetonitrile (100 mL)/water (50 mL) and lyophilized to provide **74**. LC/MS: (M+1)⁺: 846.1.

Step G: Preparation of Intermediate 75

To a solution of **74** (2.85 g, 3.23 mmol) in DMF (45 ml) was added HATU (1.474 g, 3.88 mmol), and the resulting solution was stirred at ambient temperature for 30 min, then diluted with DCM (600 ml) followed by addition of DIEA (1.692 ml, 9.69 mmol) dropwise. The resulting solution was stirred at ambient temperature for 1 hour. The solution was concentrated, and the residue was purified by reverse phase MPLC over C18 column using acetonitrile (0.05%TFA) / water (0.05%TFA) as eluting solvents. The fractions containing the product were concentrated and the aqueous layer was partitioned between DCM (200 mL) and sat. NaHCO₃ (200 mL). The aqueous phase was extracted with DCM (2x100 mL) and the combined organic phase was dried over Na₂SO₄ and concentrated to give **75**.

LC/MS: (M+1)⁺: 828.1.

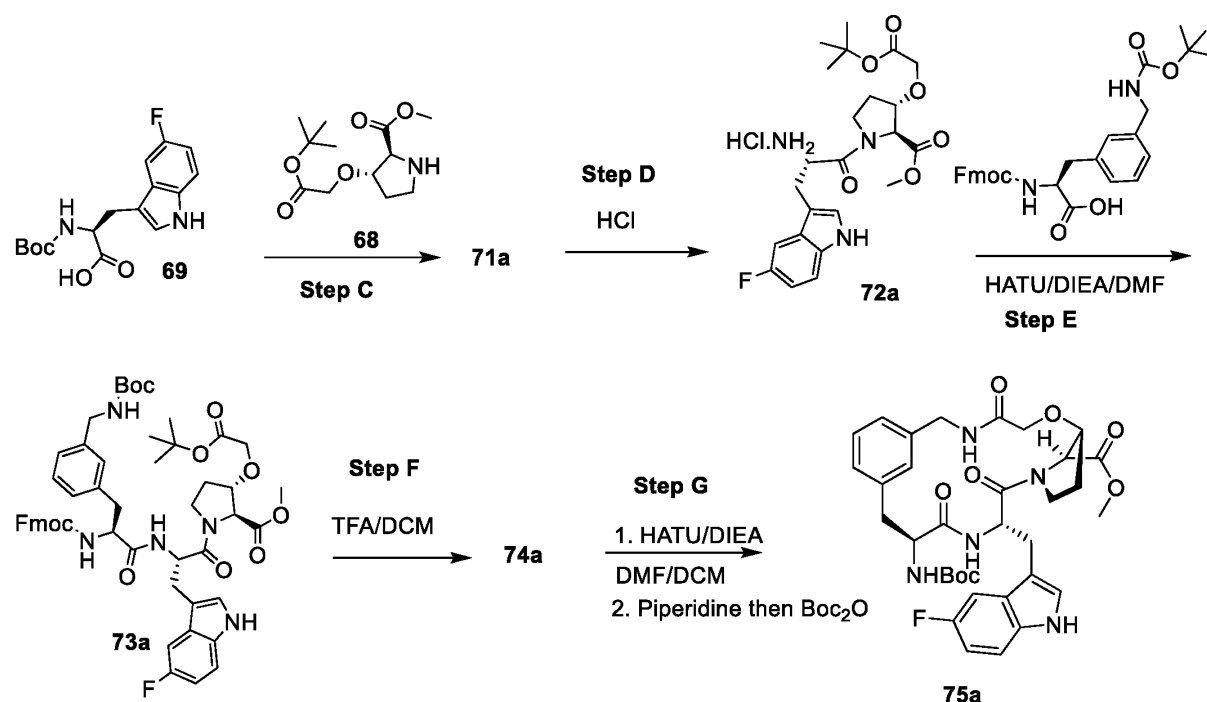
Step H: Preparation of Intermediate compound 76

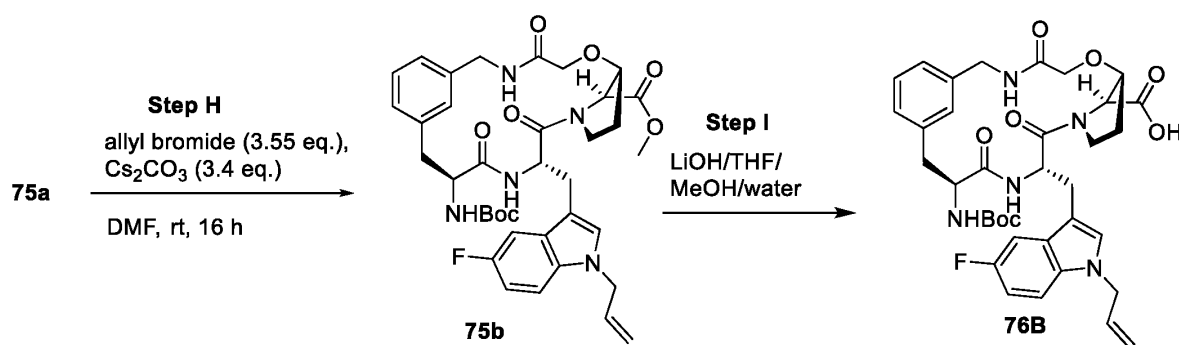
To a solution of **75** (1.93 g, 2.331 mmol) in THF (60 ml), MeOH (30 ml), and water (20 ml) at 0 °C was added 1N aqueous LiOH (9.9 ml, 9.90 mmol) dropwise, and the resulting solution was stirred at 0 °C for 16h then quenched by addition of HCl (1N, 9.9 mL). The

volatile was evaporated on rotary evaporator and to the solution above at 0 °C was added acetone (60 ml), sodium carbonate (0.371 g, 3.50 mmol), and Fmoc-Osu (0.802 g, 2.378 mmol). The resulting solution was stirred at 0 °C for 6h, the volatile was evaporated on rotary evaporator, the aqueous phase was acidified to pH 4, then extracted with

30%IPA/DCM (3x100 mL). The combined organic phase was dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using MeOH/DCM as eluting solvents to give **76**. LC/MS: (M+1)⁺: 814.2.

Alternative preparation of intermediate compound 75b and intermediate compound 76B there from:





The general procedure described above for the preparation of intermediate compound **75** was generally followed, except that Step B was eliminated and the final step G also included replacing the Fmoc protecting group with a Boc protecting group, thus providing the intermediate compound **75a**. The procedures for **75a** and **76B** are described below.

*Step C: Preparation of intermediate compound **71a***

To a solution of **69** (5.00 g, 15.5 mmol) in DMF (40.0 mL) at -50 °C were added **68**, HATU (5.90 g, 15.5 mmol) and DIEA (4.01 g, 31.0 mmol) and the reaction mixture was stirred at -50 °C for 3 h. The final solution was quenched with water (5 mL), concentrated under reduced pressure and the residue was purified by reverse phase column chromatography over C18 (eluting with a gradient of acetonitrile/water + 0.01% ammonium bicarbonate) to provide **71a**. LCMS (ESI) calc'd for C₂₈H₃₈FN₃O₈ [M + H]⁺: 564.3, found 564.2.

*Step D: Preparation of intermediate compound **72a***

To a solution of 2 N HCl in dioxane (100 mL) and THF (100 mL) at rt was added **71a** (8.40 g, 14.9 mmol) and the reaction mixture was stirred for 5 h. The final solution was concentrated under reduced pressure to afford **72a**. LCMS (ESI) calc'd for C₂₃H₃₁ClFN₃O₆ [M - HCl + H]⁺: 464.2, found 464.3.

*Step E: Preparation of intermediate compound **73a***

To a solution of **72a** (800 mg, 1.60 mmol) in DMF (10.0 mL) at -50 °C were added (S)-2-((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(3-((((tert-butoxycarbonyl)amino)methyl)phenyl)propanoic acid (827 mg, 1.60 mmol), HATU (608 mg, 1.60 mmol) and DIEA (620 mg, 4.80 mmol) and the mixture was stirred at -50 °C for 3 h.

- 5 The resulting solution was diluted with water (50 mL) and the aqueous layer was extracted with EtOAc (3 x 100 mL). The combined organic layer was washed with brine, dried over anhydrous Na₂SO₄ and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by silica gel column chromatography (eluting with a gradient 1%-60% of EtOAc in PE) to give **73a**. LCMS (ESI) calc'd for C₅₃H₆₀FN₅O₁₁ [M + H]⁺: 962.4, found 962.6.

Step F: Preparation of intermediate compound 74a

- To a solution of **73a** (3.00 g, 3.12 mmol) in DCM (15.0 mL) at rt was added TFA (15.0 mL) and the reaction mixture was stirred for 1 h at room temperature. The resulting solution was concentrated under reduced pressure, co-evaporated with toluene and DCM to give **74a**. LCMS (ESI) calc'd for C₄₆H₄₅F₄N₅O₁₁ [M - TFA + H]⁺: 806.3, found 806.7.

Step G: Preparation of intermediate compound 75a

- To a solution of **74a** (4.00 g, 4.35 mmol) in DMF (150 mL) at rt was added HATU (1.65 g, 4.35 mmol) and the reaction solution was stirred for 0.5 h. The solution was diluted with DCM (450 mL) and DIEA (1.69 g, 13.1 mmol) then stirred at rt for 3 h. The resulting solution was quenched with water (5 mL), concentrated under reduced pressure and the residue was purified by reverse phase column chromatography over C18 (eluting with a gradient of acetonitrile/water + 0.05%TFA) to provide a Fmoc-protected intermediate.

LCMS (ESI) calc'd for C₄₄H₄₂FN₅O₈ [M + H]⁺: 788.3, found 788.9.

- To a solution of the Fmoc-protected intermediate just above (200 mg, 0.250 mmol) in DCM (5.00 mL) at rt was added piperidine (1.25 mL) and the reaction solution was stirred

for 1 h. The final solution was concentrated under reduced pressure and the residue was purified by silica gel column chromatography (eluting with a gradient 0%-5% of MeOH in DCM) to afford an amine intermediate. LCMS (ESI) calc'd for $C_{29}H_{32}FN_5O_6$ $[M + H]^+$: 566.2, found 566.3.

5 To a solution of the amine intermediate just above (2.86 g, 5.06 mmol) in THF (30.0 mL) and water (30.0 mL) at rt were added Boc_2O (2.21 g, 10.1 mmol) and sodium bicarbonate (1.70 g, 20.2 mmol) and the reaction was stirred for 3 h. The final solution was diluted with water (50 mL) and extracted with EtOAc (3 x 100 mL). The combined organic layer was washed with brine (3 x 100 mL), dried over anhydrous Na_2SO_4 and filtered. The
10 filtrate was concentrated under reduced pressure and the residue was purified by silica gel column chromatography (eluting with a gradient 0%-5% MeOH in DCM) to provide **75a**.
LCMS (ESI) calc'd for $C_{34}H_{40}FN_5O_8$ $[M + H]^+$: 666.3, found 666.5; ^1H NMR (300 MHz, CD_3OD) δ 7.35-7.08 (m, 6H), 6.95-6.79 (m, 2H), 5.01-4.91 (m, 1H), 4.72-4.56 (m, 2H), 4.45-4.38 (m, 1H), 4.45-4.38 (m, 5H), 3.69 (s, 3H), 3.32-2.92 (m, 5H), 2.14-1.82 (m, 2H), 1.47 (s,
15 9H). *Step H: Preparation of intermediate compound 75b*

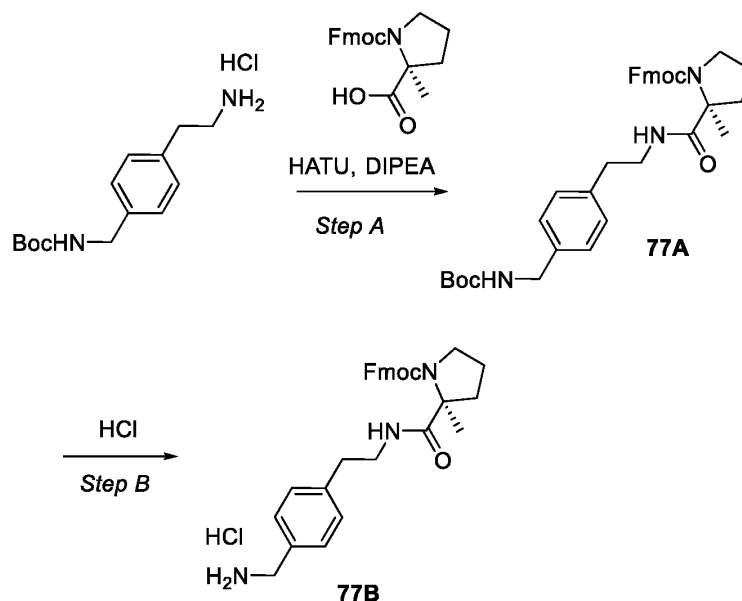
To a solution of **75a** (0.665 g, 0.99 mmol) in DMF (0.5 mL) was added Cs_2CO_3 (1.11 g, 3.40 mmol) and 3-bromoprop-1-ene (0.43 g, 3.55 mmol) at 0 °C. The reaction mixture was stirred at room temperature for 16 hours and poured into 5 mL of 50% sat. brine/10% citric acid solution, then extracted with ethyl acetate (2 x 20 mL). The organic layer was
20 washed with brine (3 x 20 mL), dried over anhydrous $MgSO_4$ and filtered. The filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography, eluting with a 1%-5% gradient of MeOH in DCM. The fractions containing intermediate compound **75b** were combined and concentrated to afford the title compound.
LCMS (ESI) calc'd for $C_{37}H_{44}FN_5O_8$ $[M + H]^+$: 706.3, found 706.3.

25 *Step I: Preparation of intermediate compound 76B*

Hydrolysis of **75b** with LiOH followed conditions similar to the ones described in the preparation of intermediate **93** to provide **76B**.

Preparation of Intermediate **77B**

5



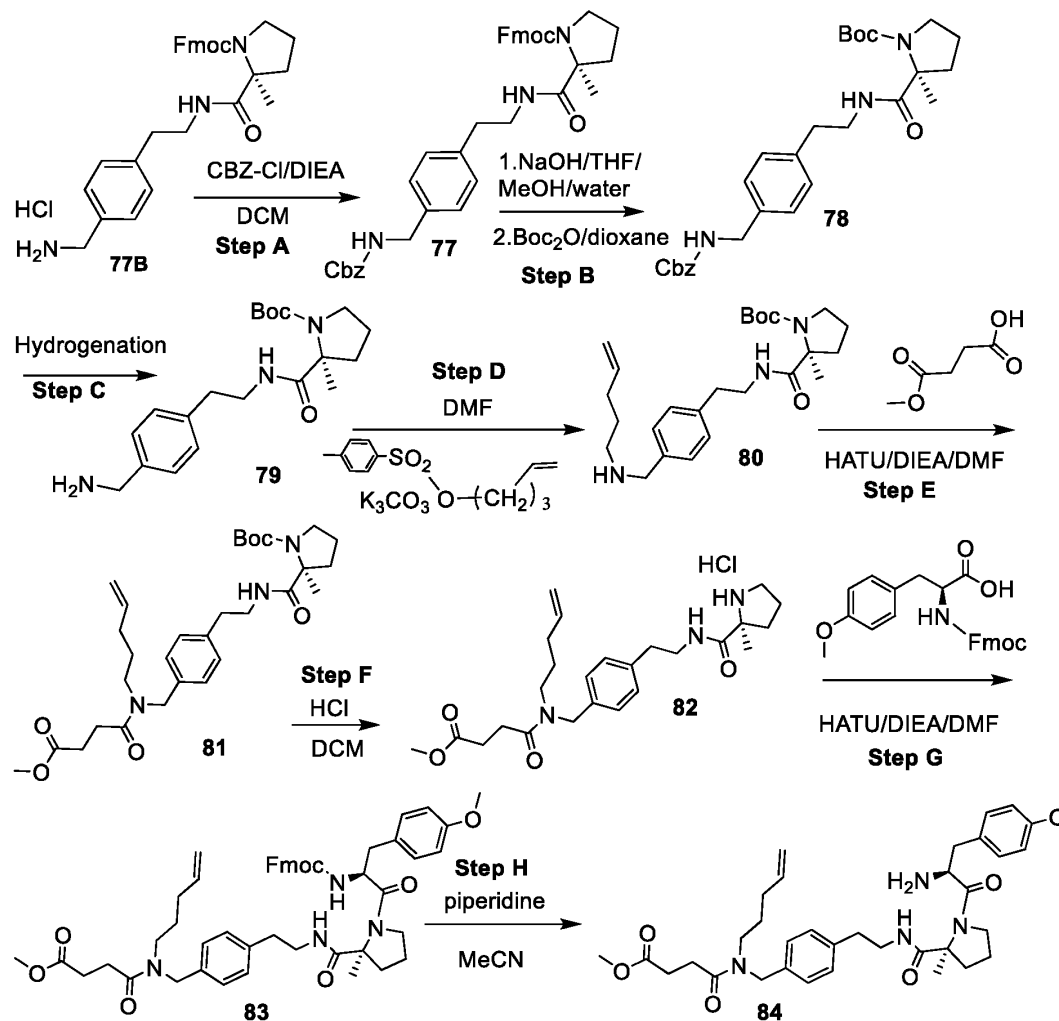
Step A: Preparation of intermediate **77A**

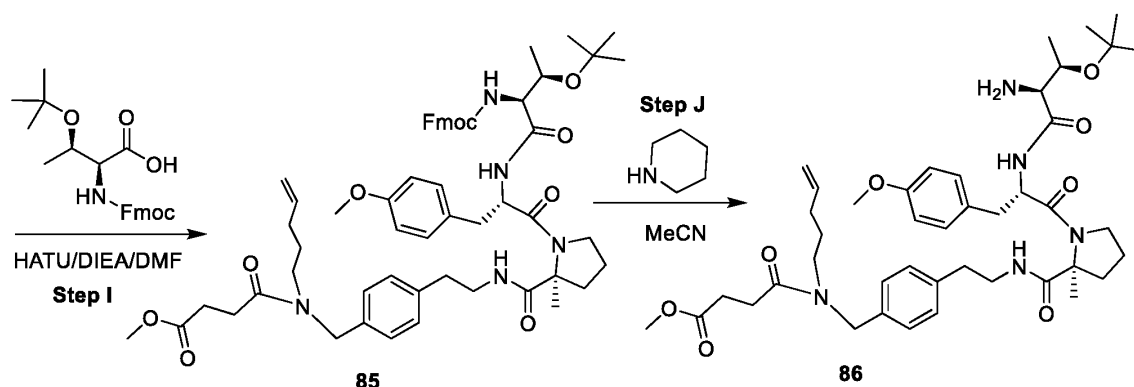
To a solution of (S)-1-(((9H-fluoren-9-yl)methoxy)carbonyl)-2-methylpyrrolidine-2-carboxylic acid (6.16 g, 17.54 mmol) and tert-butyl 4-(2-aminoethyl)benzylcarbamate hydrochloride (5.03 g, 17.54 mmol) in DMF (140 ml) at 0°C were added HATU (8.00 g, 21.05 mmol) and DIPEA (9.16 ml, 52.6 mmol) then the reaction was allowed to warm to r.t. and stirred for 2 h. The final mixture was diluted with water, extracted with EtOAc, washed with brine, dried over MgSO_4 , and filtered. The filtrate was concentrated and the residue was purified by column chromatography over silica gel (eluting with a gradient of 0-60% of EtOAc in hexane) to give **77A**. MS (ESI): m/z ($M+H$)⁺ 584.5.

Step B: Preparation of intermediate **77B**

To a solution of **77A** (8.92 g, 15.28 mmol) in DCM (40 ml) was added HCl 4N in dioxane (15.28 ml, 61.1 mmol), and the resulting solution was stirred at r.t. overnight. The mixture was concentrated to give **77B**. MS (ESI): m/z (M+H)⁺ 484.3.

5 Preparation of Intermediate 86





Step A: Preparation of intermediate 77

To a solution of (S)-(9H-fluoren-9-yl)methyl 2-((4-(aminomethyl)phenethyl)carbamoyl)-2-methylpyrrolidine-1-carboxylate hydrochloride (**77B**) (5.87 g, 11.29 mmol) in DCM (140 ml) at 0 °C was added DIEA (5.91 ml, 33.9 mmol) and CBZ-Cl (1.726 ml, 11.85 mmol) dropwise, and the resulting solution was stirred at 0 °C for 4h. The reaction solution was partitioned between water (200 mL) and DCM (200 mL), the water phase was extracted with DCM (100 mL), the combined organic phase was dried over Na₂SO₄, and the residue was purified on silica gel column using EtOAc/hexane as eluting solvents to give **77**.

LC/MS: (M+1)⁺: 618.3.

Step B: Preparation of intermediate 78

To a solution of **77** (5.56 g, 9.00 mmol) in THF (100 ml), water (50 ml), and MeOH (30 ml) was added 1N aqueous NaOH (45.0 ml, 45.0 mmol), and the resulting solution was stirred at rt for 2h. The volatile was evaporated, and to the aqueous solution was added dioxane (200 ml), and Boc₂O (2.508 ml, 10.80 mmol) in dioxane (20 mL). The resulting mixture was stirred from 0 °C to rt overnight. The volatile was evaporated on rotary evaporator, and the aqueous phase was extracted with DCM (3x150 mL). The combined organic phase was dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using EtOAc/hexane as eluting solvents to give **78**. LC/MS: (M+1)⁺: 496.2.

Step C: Preparation of intermediate 79

To a solution of **78** (4.04 g, 8.15 mmol) in MeOH (100 ml) was added 10% Pd/C (0.867 g, 0.815 mmol), and the resulting mixture was subjected to hydrogenation via H₂ balloon at rt for 1.5h. The mixture was filtered through Celite, and the filtrate was concentrated to give **79**. LC/MS: (M+1)⁺: 362.2.

5 *Step D: Preparation of intermediate 80*

To a solution of **79** (2.58 g, 7.14 mmol) in DMF (15 ml) was added pent-4-en-1-yl 4-methylbenzenesulfonate (0.858 g, 3.57 mmol) and K₂CO₃ (1.973 g, 14.27 mmol), and the resulting mixture was heated at 80 °C for 6h. After cooling down to rt, the mixture was filtered and the filtrate was purified by reverse phase MPLC using acetonitrile (0.05%TFA)/
10 water (0.05%TFA) as eluting solvents to give the product as TFA salt, which was further partitioned between DCM (100 mL) and 1N aqueous NaOH (50 mL). The aqueous phase was further extracted with DCM (2x50mL), the combined organic phase was dried over Na₂SO₄, and concentrated to give **80**. LC/MS: (M+1)⁺: 430.3.

Step E: Preparation of intermediate 81

15 To a solution of **80** (0.95 g, 2.211 mmol) in DMF (15 ml) was added 4-methoxy-4-oxobutanoic acid (0.321 g, 2.433 mmol), HATU (1.009 g, 2.65 mmol), and DIEA (0.927 ml, 5.31 mmol), and the resulting solution was stirred at rt for 1h. The solution was partitioned between EtOAc (200 mL) and brine (100 mL), the organic phase was washed with brine (2x100mL), dried over Na₂SO₄, concentrated and the residue was purified on silica gel
20 column using EtOAc/hexane as eluting solvents to give **81**. LC/MS: (M+1)⁺: 544.2.

Step F: Preparation of intermediate 82

To a solution of **81** (1.165 g, 2.143 mmol) in DCM (12 ml) was added HCl (4N in dioxane) (5.36 ml, 21.43 mmol). The resulting solution was stirred at rt for 3h, and the mixture was concentrated to give **82**. LC/MS: (M+1)⁺: 444.2.

25 *Step G: Preparation of intermediate 83*

To a solution of **82** (1.003 g, 2.089 mmol) in DMF (20 ml) was added Fmoc-L-Tyr(Me)-OH (0.959 g, 2.298 mmol), HATU (0.914 g, 2.403 mmol), and DIEA (1.095 ml, 6.27 mmol), and the resulting solution was stirred at rt for 50 min. The solution was partitioned between EtOAc (200 mL) and brine (100 mL), the organic phase was washed with brine (2x100 mL),
5 the combined organic phase was dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using EtOAc/hexane as eluting solvents to give **83**. LC/MS: (M+1)⁺: 843.4.

Step H: Preparation of intermediate 84

To a solution of **83** (1.63 g, 1.934 mmol) in acetonitrile (10 ml) was added piperidine (0.574
10 ml, 5.80 mmol), and the resulting solution was stirred at rt for 1h, then concentrated. The residue was resuspended in acetonitrile (20 mL) and concentrated again, the cycle was repeated once, and the residue was further dried under high vacuum to give **84**. LC/MS: (M+1)⁺: 621.3.

Step I: Preparation of intermediate 85

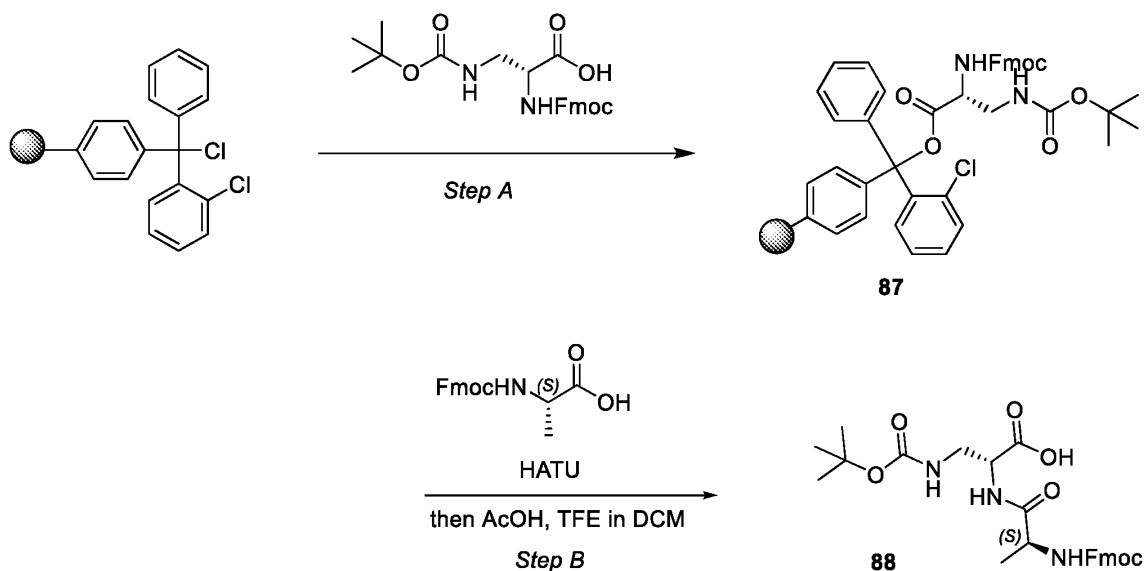
To a solution of **84** (1.2 g, 1.933 mmol) in DMF (15 ml) was added Fmoc-L-Thr(tBu)-OH (0.922 g, 2.320 mmol), HATU (0.919 g, 2.416 mmol), and DIEA (0.844 ml, 4.83 mmol), and the resulting solution was stirred at rt for 1h. The solution was partitioned between EtOAc (200 mL) and brine (100 mL), the organic phase was washed with brine (2x100 mL), dried over Na₂SO₄, concentrated and the residue was purified on silica gel column using
20 EtOAc/hexane as eluting solvents to give **85**. LC/MS: (M+1)⁺: 1000.2.

Step J: Preparation of intermediate 86

To a solution of **85** prepared in the previous step (1.94 g, 1.940 mmol) in acetonitrile (20 ml) was added piperidine (0.960 ml, 9.70 mmol), and the resulting solution was stirred at rt for 30min, then concentrated. The residue was re-dissolved in DCM/acetonitrile (1:1, 20 mL),

then concentrated again, the cycle was repeated once, and the residue was dried under high vacuum to give **86**. LC/MS: (M+1)⁺: 778.3.

Preparation of Intermediate 88



Step A – Synthesis of Intermediate 87

To 2-chloro-2-chlorotrityl resin 1-1.5 mmol/g (7.0 g, 1-1.5 mmol/g) was added dry DCM (45 ml). The resin was shaken 20 min followed by addition of half of DIPEA 0.17 N in DCM (3.67 ml, 21.00 mmol), Fmoc-D-Dap(Boc)-OH (3.28 g, 7.70 mmol) then the remainder of DIPEA 0.17 N in DCM (3.67 ml, 21.00 mmol). The resin was shaken at room temperature overnight, rinsed with DCM and dried. The resin was then quenched with 5% DIPEA and 10% MeOH in DCM (80 mL), shaken for 2 h then filtered, rinsing with DCM (3x), DMF (3x) and DCM (3x) then dried under vacuum to give resin **87** which was used as is in the next step.

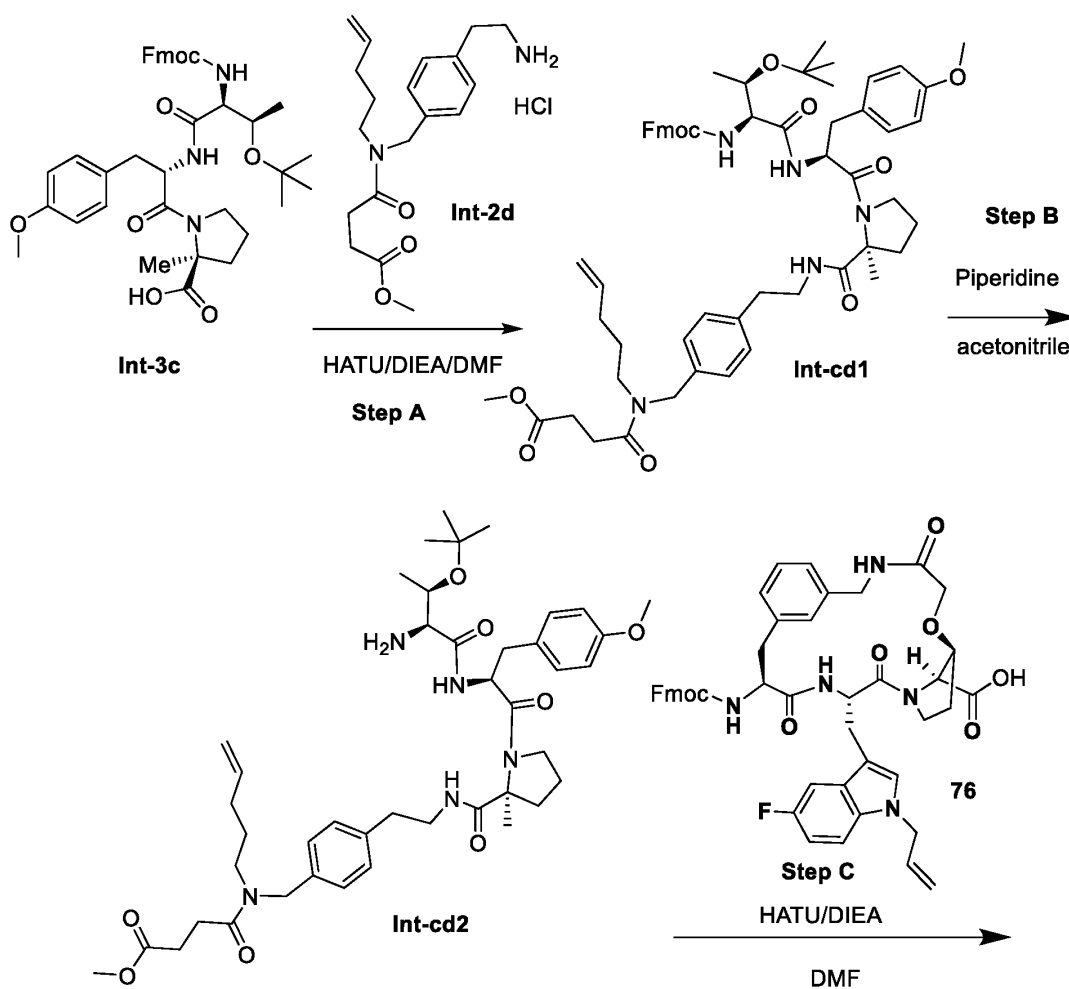
Step B – Synthesis of Intermediate 88

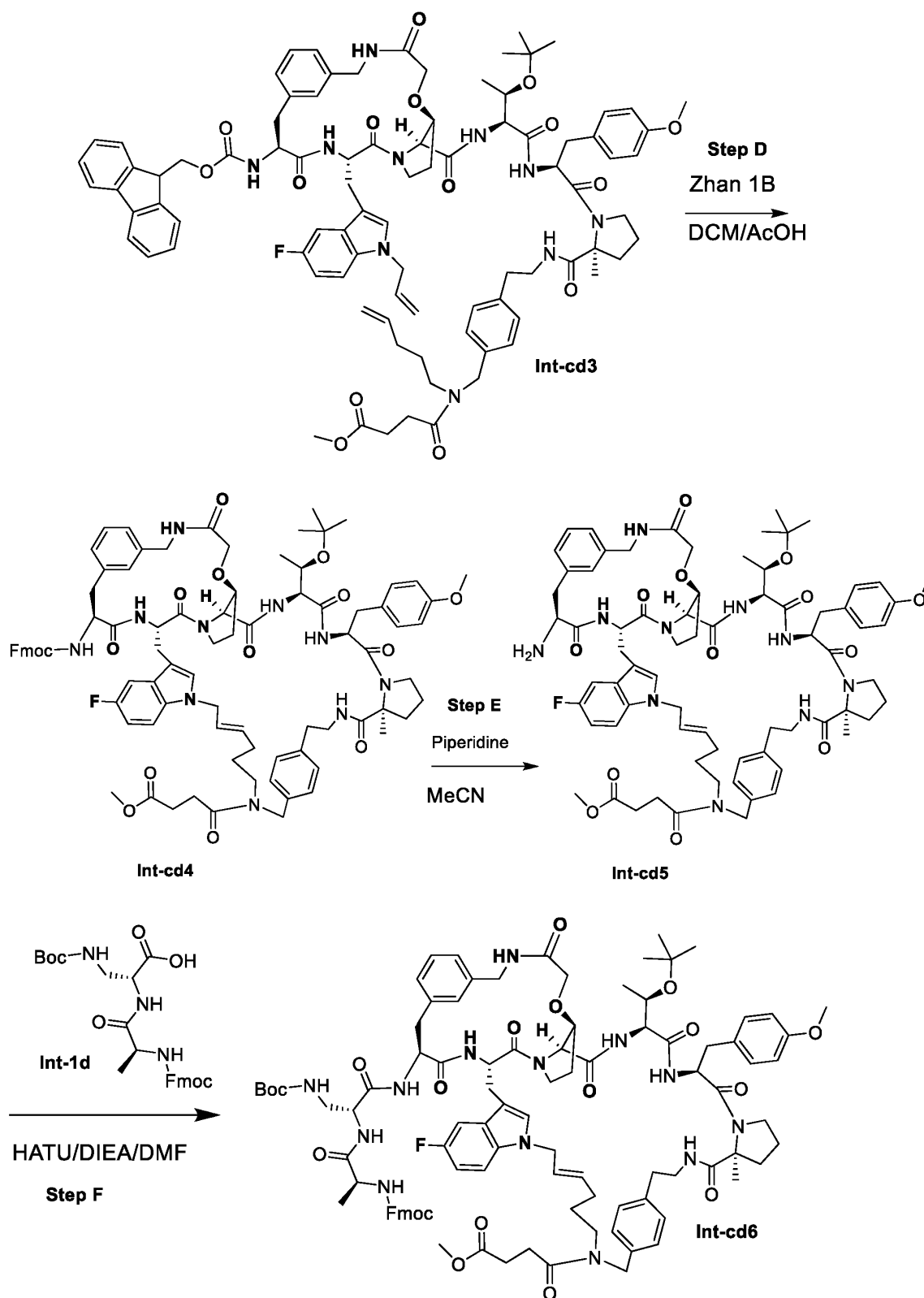
Resin **87** (4.5 g, 2.475 mmol) was manually deprotected with 5% piperidine in DMF (30 ml) for 30 min, filtered, retreated with 5% piperidine in DMF (30 ml) for another 30 min, filtered, then rinsed with DMF and DCM and dried. The resin was then manually coupled with

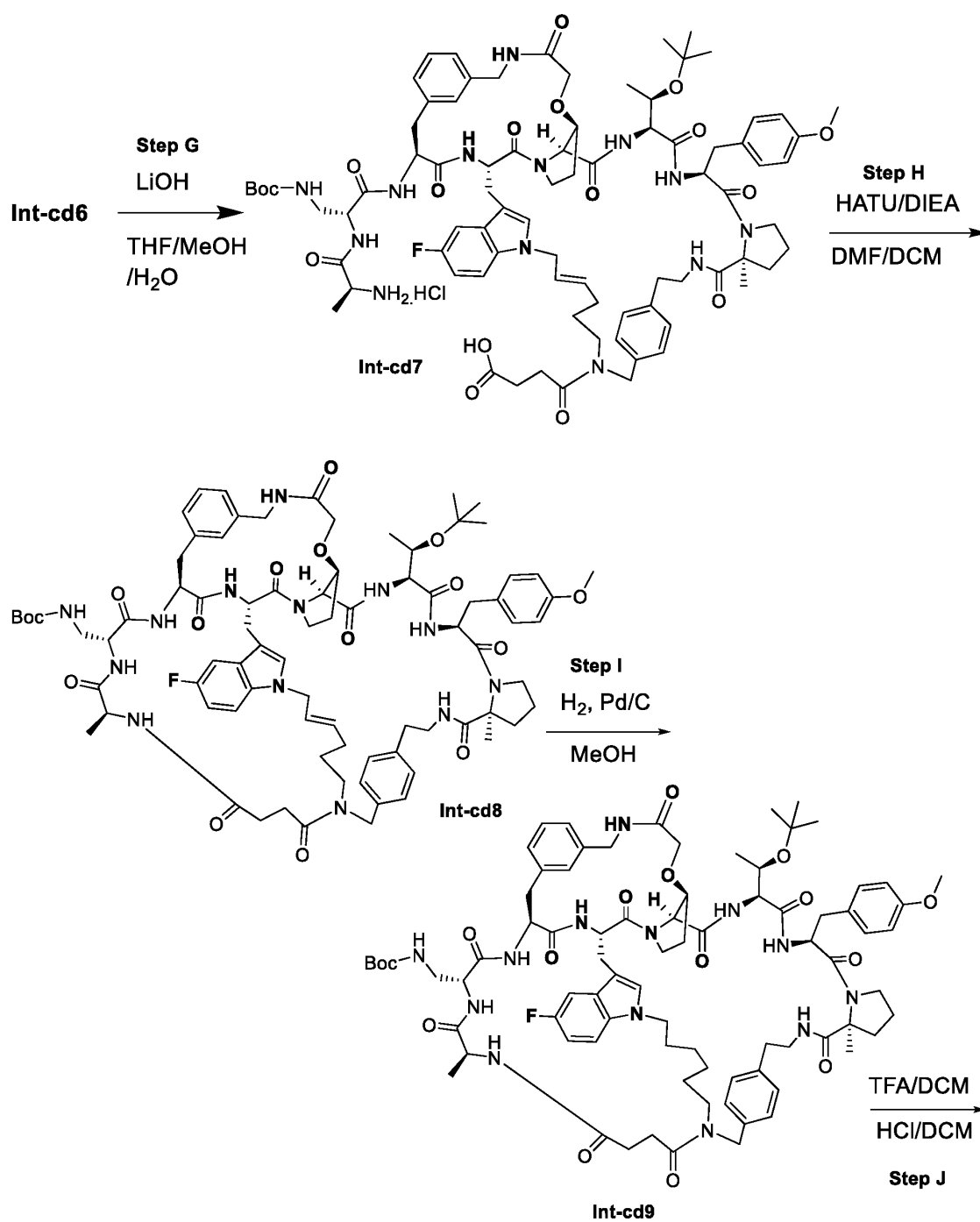
Fmoc-Ala-OH (1.541 g, 4.95 mmol), HATU (1.694 g, 4.46 mmol) and DIPEA (1.729 ml, 9.90 mmol) in DMF (30 ml) for 2 h then filtered, rinsing with DMF and DCM, then dried. The resin was then treated with 10% AcOH and TFE in DCM (60 ml) for 90 min, filtered and the filtrate was concentrated to provide **88**. LC/MS: $[2M+H]^+ = 995.01$.

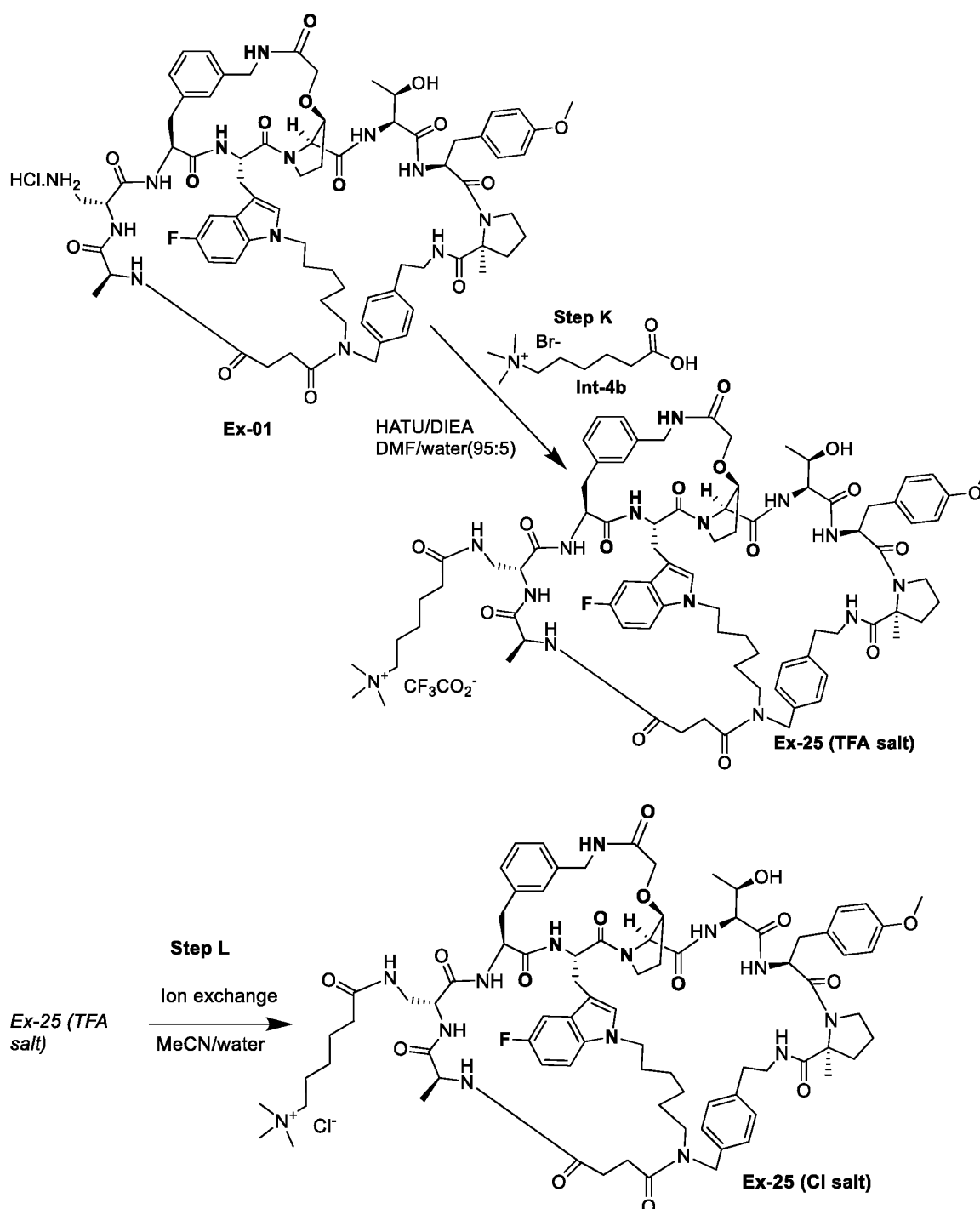
5 Example 1A – Alternative Synthesis of Ex-01 and preparation of Ex-25 therefrom:

Compound **Ex-01**, presented above, may alternatively be prepared, and compound **Ex-25** may be prepared from **Ex-01**, in accordance with the following Scheme:









Step A – Synthesis of Intermediate **Int-cd1**

To a solution of **Int-3c** (synthesis from intermediate **107** described below) (7.09 g, 10.33
 5 mmol) in DMF (45 ml) at 0 °C was added **Int-2d** (3.63 g, 9.84 mmol, preparation described

below) and HATU (3.74 g, 9.84 mmol)) followed by DIPEA in DMF (6.87 ml, 39.4 mmol) and the mixture was allowed to warm to room temperature and stirred for 1h. The mixture was quenched at 0 °C with brine and extracted with EtOAc. The combined organic fractions were washed with brine, dried over MgSO₄, filtered and concentrated in vacuo.

- 5 The residue was purified by column chromatography over silica gel (eluting with a gradient of 1% to 80% ethyl acetate in petroleum ether) to give **Int-cd1**. LC/MS: [M+1]⁺ = 1000.5.

*Step B – Synthesis of Intermediate **Int-cd2***

- To a solution of **Int-cd1** (3.48 g, 3.48 mmol) in acetonitrile (50 ml) was added piperidine (1.72 ml, 17.40 mmol), and the resulting solution was stirred at rt for 3h. The mixture was concentrated, the residue was re-dissolved in DCM/acetonitrile (1:1, 20 mL), concentrated again and the residue was dried under vacuum to give **Int-cd2** as a crude product. LC/MS: (M+1)⁺=778.5.
- 10

*Step C – Synthesis of Intermediate **Int-cd3***

- To a solution of **76** (preparation shown in Example 1, above) (2.45 g, 3.01 mmol) and **Int-cd2** (2.69 g, 3.46 mmol) in DMF (70 ml) at 0 °C was added HATU (1.37 g, 3.61 mmol) followed by DIEA (1.05 ml, 6.02 mmol). The resulting solution was stirred at rt for 50 min, then partitioned between EtOAc (500 mL) and brine (200 mL). The organic phase was washed with brine (2x200 mL), dried over Na₂SO₄, concentrated and the residue was purified by column chromatography over silica gel (eluting with a gradient of 1% – 5% MeOH in DCM) to give **Int-cd3**. LC/MS: (M+1)⁺=1574.7.
- 15
- 20

*Step D – Synthesis of Intermediate **Int-cd4***

- A room temperature solution of **Int-cd3** (1.91 g, 1.21 mmol) in DCM (1500 ml) and acetic acid (30 mL) was bubbled with N₂ for 30 min followed by addition of Zhan's catalyst-1B (0.445 g, 0.607 mmol). The resulting mixture was further bubbled at room temperature with N₂ for 30 min, then heated at 55 °C for 5h. After cooling to rt the mixture was filtered over
- 25

Celite, the filtrate was concentrated, and the residue was purified by column chromatography over silica gel (eluting with a gradient of 1% – 5% MeOH in DCM) to give **Int-cd4** (as mixture of cis and trans olefins). LC/MS: (M+1)⁺=1546.8.

*Step E – Synthesis of Intermediate **Int-cd5***

- 5 To a solution of **Int-cd4** (mixture of cis and trans olefins) (5.49 g, 3.55 mmol) in DCM (20 ml) and acetonitrile (50 ml) was added piperidine (1.76 ml, 17.8 mmol). The resulting solution was stirred at rt for 2h, then concentrated and the residue was suspended in acetonitrile (20 ml) and concentrated again. The residue was then dried under vacuum to give **Int-cd5** (as mixture of cis and trans olefins) as a crude mixture. LC/MS:
- 10 (M+1)⁺=1323.8.

*Step F – Synthesis of Intermediate **Int-cd6***

- To a solution of **Int-cd5** (mixture of cis and trans olefins) (4.70 g, 3.55 mmol) and **Int-1d** (2.21 g, 4.44 mmol, preparation described below) in DMF (70 ml) at 0 °C was added HATU (1.76 g, 4.62 mmol) and DIEA (1.55 ml, 8.88 mmol). The resulting solution was warmed to
- 15 room temperature and stirred for 1h, then partitioned between EtOAc (300 mL) and brine (200 mL). The aqueous phase was extracted with EtOAc (200 mL), the EtOAc phase was combined and washed with brine (3x200 mL), dried over Na₂SO₄, concentrated and the residue was purified by column chromatography over silica gel (eluting with a gradient of 1% – 5% MeOH in DCM) to give **Int-cd6** (as mixture of cis and trans olefins). LC/MS:
- 20 (M+1)⁺= 1802.8

*Step G – Synthesis of Intermediate **Int-cd7***

- To a solution of **Int-cd6** (as mixture of cis and trans olefins) (5.41 g, 3.00 mmol) in THF (100 ml), MeOH (30 ml), and water (30 ml) at 0 °C was added 1N aqueous LiOH (24.0 ml, 24.0 mmol) dropwise, and the resulting solution was stirred at 0 °C for 3h. The mixture was
- 25 neutralized at 0 °C by addition of 1N HCl, the volatile was evaporated, and the aqueous

layer was neutralized to pH 5 by 1N HCl. The mixture was then frozen and lyophilized, and the residue was purified by column chromatography over C18 (eluting with a gradient of acetonitrile (0.05%TFA)/water (0.05%TFA)) to give **Int-cd7** (as mixture of cis and trans olefins) as a TFA salt. To the **Int-cd7** TFA salt thus obtained, (as mixture of cis and trans olefins) in acetonitrile (750 mL) and water (450 mL) was added at 0 °C 0.1N aqueous HCl (150 ml, 15.00 mmol) dropwise, then the resulting solution was stirred at 0 °C for 5 min, frozen and lyophilized to give **Int-cd7** as a HCl salt (as mixture of cis and trans olefins). LC/MS: (M+1)⁺=1566.6.

*Step H – Synthesis of Intermediate **Int-cd8***

To a solution of **Int-cd7** HCl salt obtained from the previous step (1.01 g, 0.630 mmol) in DMF (50 ml) and DCM (1300 ml) was added DIEA (0.330 ml, 1.890 mmol) and HATU (0.287 g, 0.756 mmol). The resulting solution was stirred at rt for 2h, the volatile was evaporated, and the residue was partitioned between EtOAc (400 mL) and brine (200 mL). The aqueous phase was extracted with EtOAc (300 mL), the combined organic layers were washed with brine (3x100 mL), dried over Na₂SO₄, concentrated and the residue was purified by column chromatography over silica gel (eluting with a gradient of 1% – 10% MeOH in DCM) to give **Int-cd8** (as mixture of cis and trans olefins). LC/MS: (M+1)⁺=1548.8.

*Step I – Synthesis of Intermediate **Int-cd9***

To a solution of **Int-cd8** obtained in the previous step (1.22 g, 0.788 mmol) in MeOH (100 ml) was added 10% Pd/C (0.645 g, 0.607 mmol), and the resulting mixture was hydrogenated at ambient temperature via H₂ balloon for 7 hours. After 7 hours the reaction was filtered over Celite, the filtrate was concentrated, and the residue was purified by column chromatography over silica gel (eluting with a gradient of 1% – 10% MeOH in DCM) to give **Int-cd9**. LC/MS: (M+1)⁺=1550.9.

*Step J – Synthesis of compound **Ex-01** as the -HCl salt*

To a solution of **Int-cd9** (1.14 g, 0.735 mmol) in DCM (6 ml) was added TFA (12 ml, 156 mmol), and the resulting solution was stirred at ambient temperature for 30 min. The mixture was then concentrated, and the residue was dissolved in DCM (20 mL) and toluene
5 (20 mL). The resulting mixture was concentrated, and the residue was re-dissolved in DCM (20 mL) and treated with HCl (4N in dioxane) (0.919 ml, 3.68 mmol). The resulting mixture was concentrated to give the product as solid. This solid product was re-dissolved in acetonitrile (200 mL) and water (100 mL), and to the above solution at 0 °C was added 1N aqueous HCl (3.68 ml, 3.68 mmol) dropwise. The resulting solution was stirred at 0 °C for 2
10 min, then frozen and lyophilized to give **Ex-01** as a -HCl salt. LC/MS: (M+1)⁺=1394.7.

*Step K – Synthesis of Example **Ex-25** as the TFA salt*

To a solution of **Ex-01** HCl salt (870 mg, 0.608 mmol) and **Int-4b** (170 mg, 0.669 mmol, preparation described below) in DMF (1.2 ml) and water (0.6 ml) was added HATU (254 mg, 0.669 mmol) and DIEA (425 µl, 2.433 mmol). The resulting solution was stirred at rt for
15 1h then quenched by addition of 1.2 mL water. The mixture was filtered, and the filtrate was purified by column chromatography over C18 (eluting with a gradient of acetonitrile (0.05%TFA)/water (0.05%TFA)) to give **Ex-25** as a TFA salt. LC/MS: M⁺= 1550.6.

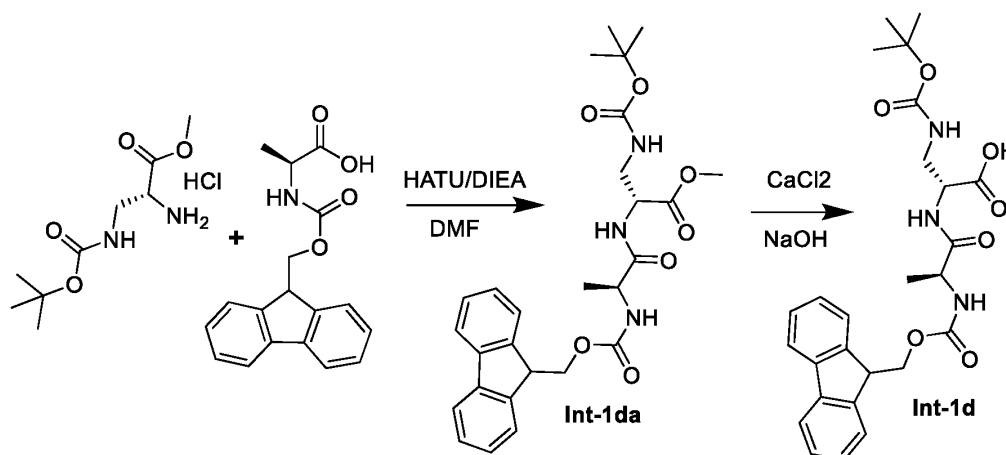
*Step L – Preparation of Example **Ex-25** as the Cl salt*

Into two columns was packed 73.6 g of AG MP-1 ion exchange resin chloride form (cat#
20 141-1841 BIO-RAD) for a total of 36.8 g resin in each column. Each column was washed with water (2x80 ml), followed by 20% acetonitrile in water (2x100ml). A solution of **Ex-25** TFA salt prepared in the previous step (737 mg, 0.443 mmol) in 20% acetonitrile in water (100 mL) was loaded evenly onto the two resin columns, then each column was eluted with 20% acetonitrile in water (130 ml). The eluents were combined, frozen and lyophilized to
25 give **Ex-25** as the chloride salt. LC/MS: M⁺ = 1550.6.

There follows the description of a number of intermediates which are usefully employed in the synthesis of **Ex-01** and **Ex-25** described immediately preceding.

Preparation of Intermediate Int-1d

Intermediate compound **Int-1d** was prepared from starting materials in accordance with the following scheme:



Step A – Synthesis of Int-1da

To a solution of D-Dap(Boc)-OMe HCl salt (4.10 g, 16.10 mmol), Fmoc-Ala-OH (5.01 g, 16.10 mmol) and HATU (6.43 g, 16.90 mmol) in DMF (40 ml) at 0 °C was added DIPEA (7.03 ml, 40.2 mmol) and the mixture was stirred at 0 °C for 2 h then kept in the refrigerator overnight. The mixture was quenched at room temperature with water and extracted with EtOAc. The combined organic fractions were washed with half brine, dried over Na_2SO_4 , filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with a gradient of Hexanes/EtOAc) to give **Int-1da**.

LC/MS: $[\text{M}+\text{H}]^+ = 512.3$.

Step B – Synthesis of Int-1d

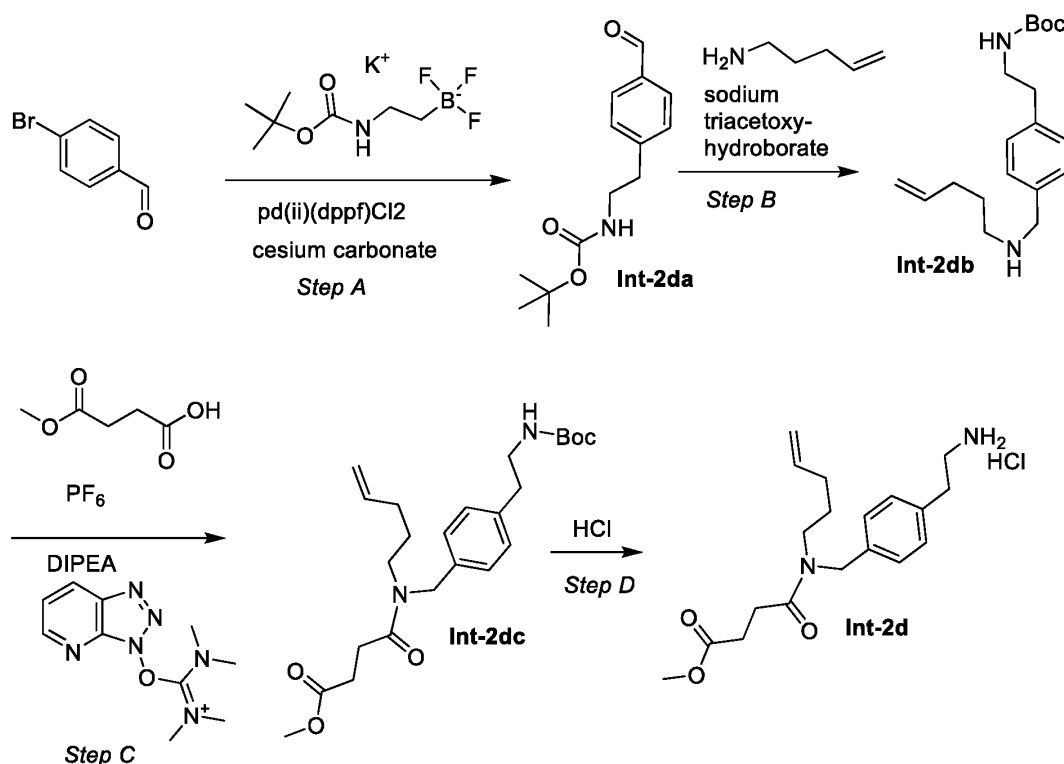
To a solution of **Int-1da** (8.03 g, 15.70 mmol) and 0.8 N calcium chloride (19.62 ml, 15.70 mmol) in water (40 ml) and 2-propanol (120 ml) at room temperature was added solid sodium hydroxide (0.691 g, 17.27 mmol). The mixture was stirred at room temperature

overnight. The mixture was concentrated, acidified with 0.5 N to pH ~2 (~40 mL), extracted trice with EtOAc, washed with brine, dried over Na₂SO₄ and concentrated. The residue was purified by column chromatography over C18 (eluting with a gradient of acetonitrile/water + 0.1% TFA) to give **Int-1d**. LC/MS: [M+H]⁺ = 498.25.

5 The preparation of intermediate **Int-1d** is described above for use in the preparation of **Ex-01** and **Ex-25**. This portion of the molecule may be described as a “linker” which cyclizes the lower peptide ring to the higher peptide ring. Other similar “linkers” may be used in place of **Int-1d** by varying the spacer used in the synthesis, including but not limited to, using Dap and D-Ala.

10 Preparation of intermediate Int-2d

Intermediate **Int-2d**, useful as a “linker” in the preparation of compounds of the invention, was prepared in accordance with the following scheme:



15 Step A – Synthesis of Int-2da

A solution of 4-bromobenzaldehyde (15.00 g, 81 mmol), potassium tert-butyl N-[2-(trifluoroboranuidyl)ethyl]carbamate (20.97 g, 84 mmol), cesium carbonate (52.8 g, 162 mmol) and 1,1'-bis(diphenylphosphino)ferrocene-palladium(ii)dichloride dichloromethane complex (Pd(II)(dppf)Cl₂, 1.99 g, 2.43 mmol) in degassed toluene (250 ml) and water (85
5 ml) was warmed to 76 °C and stirred overnight. The mixture was quenched at room temperature with half-saturated aqueous ammonium chloride and extracted with EtOAc. The combined organic fractions were washed with brine, dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with a gradient of DCM/EtOAc) to give **Int-2da**. LC/MS: (M-56+1)⁺= 193.0.

10 *Step B – Synthesis of Int-2db*

To a solution of **Int-2da** (12.9 g, 51.7 mmol) and pent-4-en-1-amine (6.61 g, 78 mmol) in DCM (120 ml) and AcOH (3 ml) at room temperature in a water bath was added sodium triacetoxhydroborate (32.9 g, 155 mmol) portion wise and the mixture was stirred for 30 min. The reaction was slowly quenched at 0 °C with 3 ml of water, poured into 1 N NaOH
15 (500 ml), stirred for 15 min then extracted with DCM, dried over Na₂SO₄, and concentrated. The residue was purified by column chromatography over silica gel (eluting with a gradient of DCM/MeOH) to give **Int-2db**. LC/MS: (M+1)⁺= 319.2.

Step C – Synthesis of Int-2dc

To a solution of **Int-2db** (8.48 g, 20.77 mmol) and 4-methoxy-4-oxobutanoic acid (3.02 g, 22.85 mmol) in DMF (40 ml) was added HATU (9.48 g, 24.92 mmol) and DIPEA (8.71 ml, 49.8 mmol). The resulting solution was stirred at room temperature for 1 hour, then
20 quenched with aqueous saturated NaHCO₃ (10 mL). The mixture was partitioned between EtOAc (500 mL) and aqueous saturated NaHCO₃ (200 mL), the organic phase was washed with brine (3x200 mL), dried over Na₂SO₄, concentrated and the residue was purified on

silica gel column (eluting with a gradient of Hexanes/EtOAc) to give **Int-2dc**. LC/MS:

$(M+1)^+ = 433.4$.

Step D – Synthesis of **Int-2d**

To a solution of **Int-2dc** (2.9 g, 6.70 mmol) in DCM (15 mL) was added 4 M HCl in Dioxane

5 (10 mL) at room temperature. The reaction mixture was stirred at room temperature for 1 h.

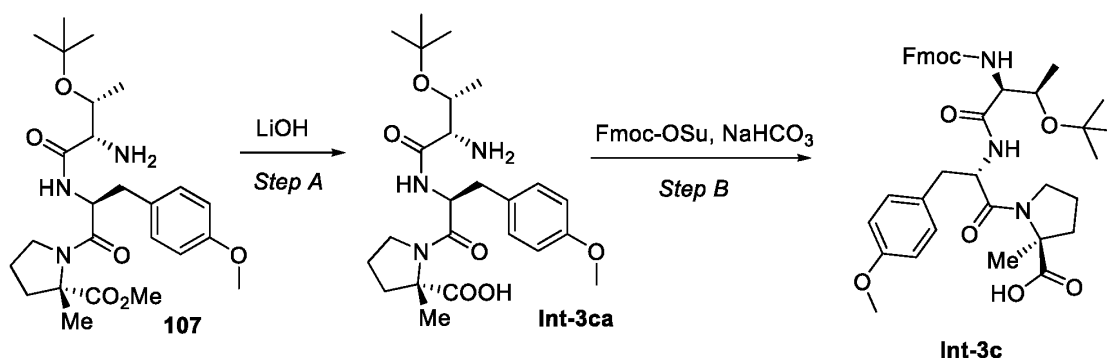
The mixture was concentrated under reduced pressure to afford methyl 4-((4-(2-

aminoethyl)benzyl)(pent-4-en-1-yl)amino)-4-oxobutanoate hydrochloride (**Int-2d**). LC/MS

$[M-HCl + H]^+ = 333.3$.

Preparation of **Int-3c** from **107** used in the synthesis of **Ex-01** and **Ex-25** described above

10 Intermediate **Int-3c** was prepared in accordance with the following scheme:



Step A – Synthesis of **Int-3ca**

Into a solution of **107** (the preparation of which is presented in the synthesis of **109** and

15 later used for **116**, which intermediate compound is used in the synthesis of Ex-53, Ex-54

and Ex-55 in Example 3 below) (10.34 g, 21.65 mmol) in THF (100 ml) at room

temperature, was added 2 N lithium hydroxide monohydrate (43.3 ml, 87 mmol) and the

mixture was warmed to 45 °C and stirred overnight to give **Int-3ca** as crude solution.

LC/MS: $(M+1)^+ = 464.3$. The reaction mixture was cooled to 0 °C and treated with 1 M HCl

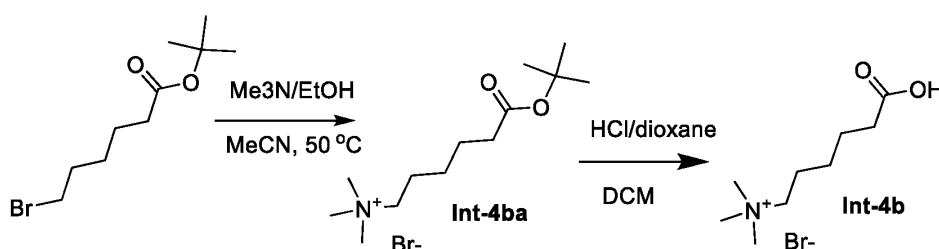
20 (40 mL). The mixture was directly used for the next step.

Step B – Synthesis of Int-3c

To crude **Int-3ca** prepared in the previous step was added NaHCO_3 (1.725 g, 20.54 mmol) and Fmoc-OSu (3.81 g, 11.30 mmol) at 0 °C. The reaction mixture was stirred at 0 °C for 2 h, treated with 1 M HCl (20.5 mL), and extracted with ethyl acetate (2 x 200 mL). The combined organic layers were washed with brine (2 x 100 mL) and dried over anhydrous sodium sulfate, filtered and concentrated under reduced pressure. The residue was purified by silica gel column chromatography, eluting with a gradient of 2% to 5% MeOH in DCM, to afford **Int-3c**. LC/MS: $(\text{M}+1)^+ = 686.4$.

Preparation of intermediate Int-4b

Intermediate **Int-4b** was prepared in accordance with the following scheme:

*Step A- Synthesis of Int-4ba from tert-butyl 3-(2-hydroxyethoxy)propanoate*

To a solution of *tert*-butyl 3-(2-hydroxyethoxy)propanoate (500.0 mg, 2.63 mmol) in DCM (2 mL) were added CBr_4 (1395 mg, 4.21 mmol) and PPh_3 (965 mg, 3.68 mmol) at 0 °C. The mixture was stirred at room temperature for 2 h. The resulting mixture was concentrated under reduced pressure and the residue was purified by silica gel column chromatography, eluting with a gradient 1% - 15% of ethyl acetate in petroleum ether. The fractions containing the desired product were combined and concentrated to afford *tert*-butyl 6-bromohexanoate. Thus prepared, a solution of *tert*-butyl 6-bromohexanoate (5 g, 19.91 mmol) in acetonitrile (10 mL) was treated with trimethylamine (13.56 mL, 59.7 mmol) and the resulting solution was heated at 50°C overnight. The solution was concentrated to give **Int-4ba**. LC/MS: $\text{M}^+ = 230.3$.

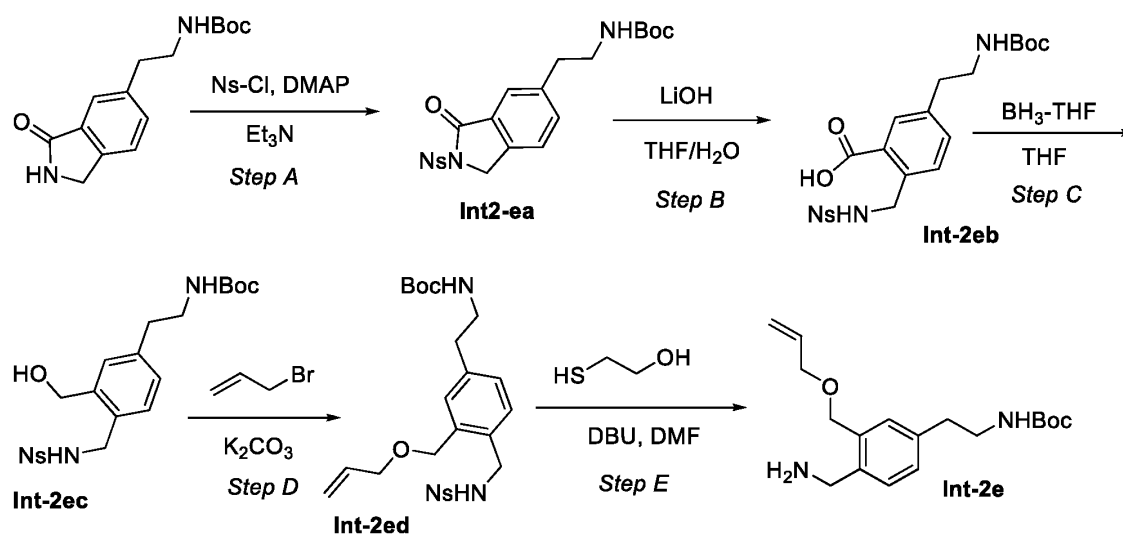
Step B – Synthesis of Int-4b

To a solution of **Int-4ba** (6.8 g, 21.92 mmol) in DCM (6 ml) was added 4N HCl in dioxane (27.4 ml, 110 mmol), and the resulting solution was stirred at rt for 3h. The mixture was then concentrated to give **Int-4b**. LC/MS: $M^+ = 174.3$.

- 5 The preparation of intermediate **Int-2d** is described above for use in the preparation of **Ex-01** and **Ex-25**. This portion of the molecule may be described as a “linker” which cyclizes the lower peptide ring bearing the R^1 , R^2 and R^8 substituents. Other similar “linkers” may be used in place of **Int-2d**. Following is a description of other “linkers” which may be used to prepare examples of the invention described herein.

10 **Preparation of intermediate Int-2e**

Intermediate **Int-2e**, useful as a “linker” in the preparation of compounds of the invention, was prepared in accordance with the following scheme:

15 *Step A – Synthesis of Intermediate Int-2ea*

To a solution of tert-butyl (2-(3-oxoisindolin-5-yl)ethyl)carbamate (1.60 g, 5.79 mmol) in DCE (20 mL) were added NsCl (1.93 g, 8.69 mmol), triethylamine (1.76 g, 17.4 mmol) and DMAP (0.141 g, 1.16 mmol). The reaction mixture was stirred for 14 h at 40 °C. The reaction was cooled to room temperature and concentrated under reduced pressure. The

residue was purified by column chromatography over silica gel (eluting with a 1%-40% gradient of EtOAc in PE) to give **Int-2ea**. LC/MS: (M+Na)⁺= 484.4.

Step B – Synthesis of Intermediate Int-2eb

To a solution of **Int-2ea** (11.3 g, 24.5 mmol) in THF (100 mL) and water (100 mL) was
5 added LiOH (1.76 g, 73.5 mmol). The reaction mixture was stirred for 5 h at 25 °C then the
resulting solution was adjusted to pH 4~5 with HCl (1M). The solution was extracted with
EtOAc and the combined organic layers were washed with brine, dried over anhydrous
Na₂SO₄ and filtered. The filtrate was concentrated under reduced pressure and the
residue was purified by column chromatography over silica gel (eluting with a 1%-6%
10 gradient of MeOH in DCM) to afford **Int-2eb**. LC/MS: (M+Na)⁺= 502.2.

Step C – Synthesis of Intermediate Int-2ec

To a solution of **Int-2eb** (1.70 g, 3.55 mmol) in THF (8 mL) was added borane (0.147 g,
10.6 mmol) at 0 °C. The reaction mixture was stirred for 14 h at 25 °C then the resulting
solution was concentrated under reduced pressure. The residue was purified by column
15 chromatography over silica gel (eluting with a 1%-50% gradient of EtOAc in PE) to give **Int-2ec**. LC/MS: (M+NH₄)⁺= 483.2.

Step D – Synthesis of Intermediate Int-2ed

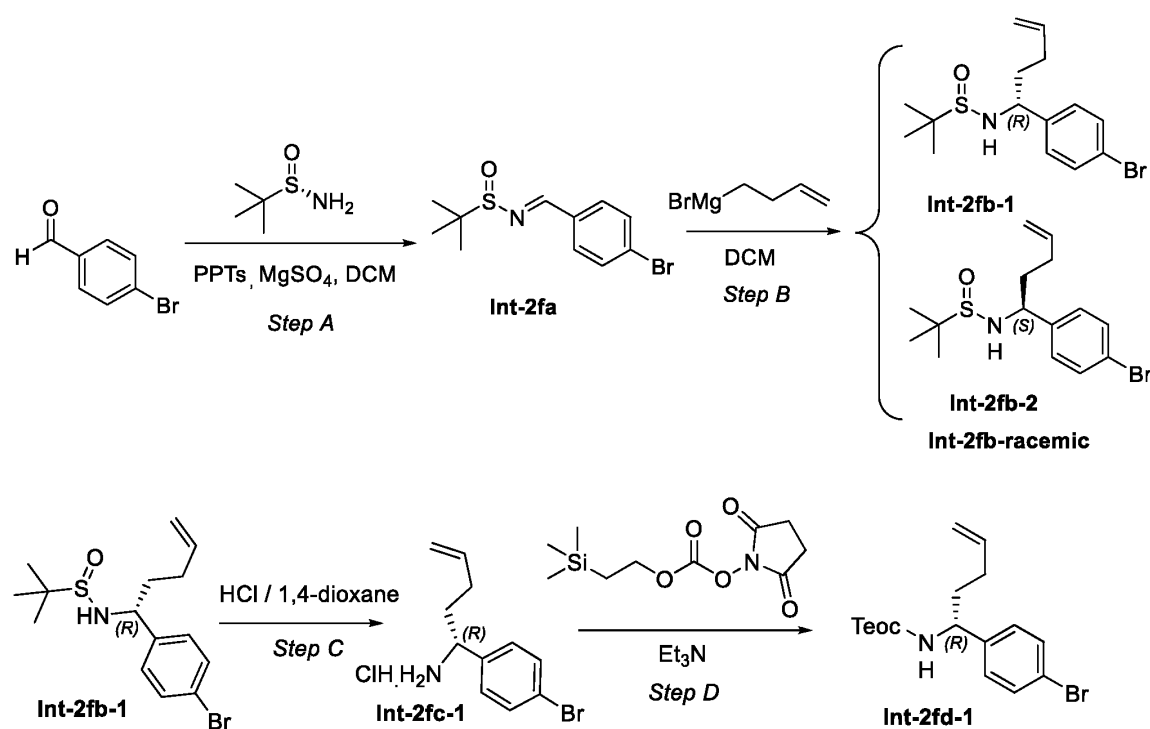
To a solution of **Int-2ec** (4.50 g, 9.67 mmol) in DMF (150 mL) was added K₂CO₃ (2.01 g,
14.5 mmol) and 3-bromoprop-1-ene (1.41 g, 11.6 mmol). The reaction mixture was stirred
20 for 5 h at room temperature then diluted with water and extracted with EtOAc. The
combined organic layers were washed with brine, dried over anhydrous Na₂SO₄ and
filtered. The filtrate was concentrated under reduced pressure and the residue was purified
by column chromatography over silica gel (eluting with a 1%-50% gradient of EtOAc in PE)
to afford **Int-2ed**. LC/MS: (M+H)⁺= 506.2.

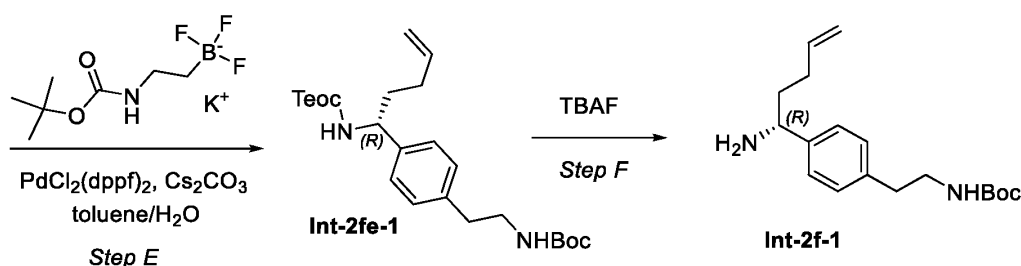
25 *Step E – Synthesis of Intermediate Int-2e*

To a solution of **Int-2ed** (4.50 g, 8.90 mmol) in DMF (35 mL) was added DBU (1.35 g, 8.90 mmol) and 2-mercaptoethanol (2.08 g, 26.7 mmol). The reaction mixture was stirred for 14 h at room temperature then purified by column chromatography over C18 (Column: 330 g; Mobile Phase A: water/0.05% TFA, Mobile Phase B: ACN; Flow rate: 85 mL/min; Gradient: 10% B to 20% B in 15 min, 20% B to 45% B in 15 min Detector: UV 210 nm; Rt=20 min) to afford **Int-2e**. LC/MS: (M+H)⁺ = 321.2. ¹H NMR (300 MHz, CDCl₃) δ 7.21-7.13 (m, 2H), 7.10-7.01 (m, 1H), 5.96-5.79 (m, 1H), 5.30-5.09 (m, 2H), 4.58 (s, 2H), 3.84 (s, 2H), 3.43-3.19 (m, 4H), 2.76 (t, J = 7.1 Hz, 2H), 1.41 (s, 9H).

Preparation of intermediate Int-2f-1

Intermediate **Int-2f-1**, useful as a "linker" in the preparation of compounds of the invention, was prepared in accordance with the following scheme:





Step A – Synthesis of Intermediate **Int-2fa**

To a solution of 4-bromobenzaldehyde (20.0 g, 108 mmol), (S)-2-methylpropane-2-sulfonamide (12.5 g, 103 mmol), MgSO_4 (130 g, 1081 mmol) in DCM (225 mL) was added
 5 pyridine 4-methylbenzenesulfonate (1.35 g, 5.40 mmol) under nitrogen protection. This mixture was stirred at 25 °C for 72 h then the resulting solution was filtered, and the filtrate was concentrated under reduced pressure. The resulting residue was purified by column chromatography over silica gel (eluting with a 1%-15% gradient of EtOAc in PE) to give **Int-2fa**. LC/MS: $(\text{M}+\text{H})^+ = 287.9, 289.9$.

10 Step B – Synthesis of Intermediate **Int-2fb** (racemate) and separation into enantiomers **Int-2fb-1** and **Int-2fb-2**

To a solution of **Int-2fa** (20.0 g, 65.9 mmol) in dry DCM (200 mL) was added but-3-en-1-ylmagnesium bromide (15.7 g, 99 mmol) slowly at -48 °C under nitrogen protection. The mixture was stirred at -48 °C for 2 h then quenched with saturated NH_4Cl (400 mL) aqueous
 15 solution and extracted with DCM. The combined organic layers were washed with brine, dried over anhydrous Na_2SO_4 and filtered. The filtrate was concentrated under reduced pressure and the resulting residue (containing **Int-2fb** racemic mixture) was purified by column chromatography over silica gel (eluting with a 1%-35% gradient of EtOAc in PE) to give **Int-2fb-1** and **Int-2fb-2**. LC/MS: $(\text{M}+\text{H})^+ = 344.0, 346.0$.

20 Step C – Synthesis of Intermediate **Int-2fc-1**

To a solution of HCl (100 mL, 4 N in 1,4-dioxane) at room temperature was added **Int-2fb-1** (17.0 g, 46.9 mmol). The reaction solution was stirred for 1 h then concentrated under reduced pressure to afford **Int-2fc-1**. LC/MS: (M+H-HCl)⁺= 240.0, 242.0.

Step D – Synthesis of Intermediate Int-2fd-1

- 5 To a solution of **Int-2fc-1** (8.20 g, 28.2 mmol) and Teoc-OSu (8.03 g, 31.0 mmol) in 1,4-dioxane (200 mL) was added TEA (8.55 g, 84 mmol) at 25 °C. This mixture was stirred for 2 hours then quenched with water and extracted with petroleum ether (PE). The combined organic layers were concentrated under reduced pressure and the residue was purified by column chromatography over silica gel (eluting with a 1%-10% gradient of EtOAc in PE) to
- 10 give **Int-2fd-1**. LC/MS: (M+Na+CH₃CN)⁺= 447.3, 449.3.

Step E – Synthesis of Intermediate Int-2fe-1

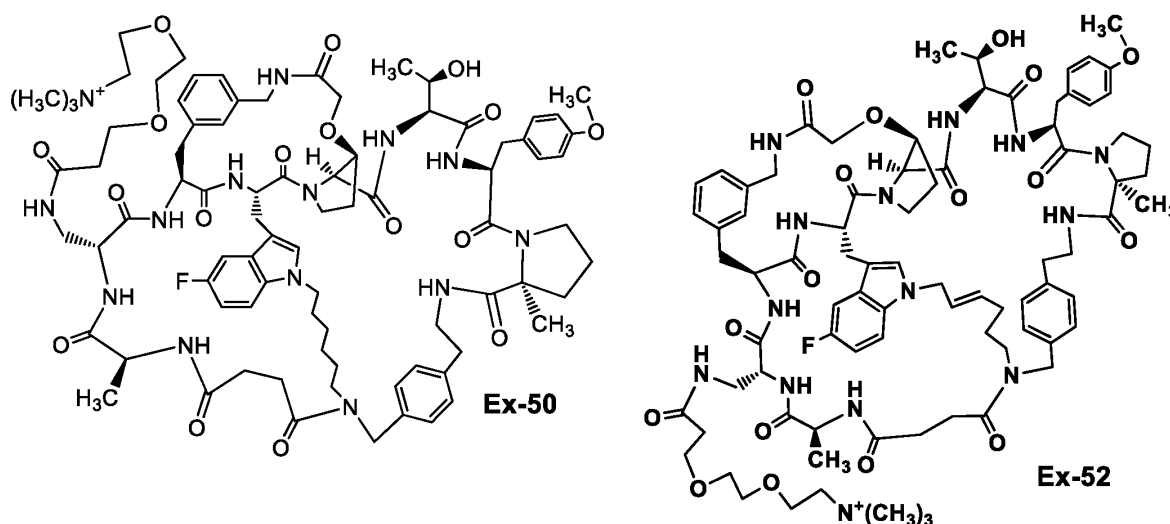
- To a solution of **Int-2fd-1** (15.1 g, 37.3 mmol), potassium (2-((tert-butoxycarbonyl)amino)ethyl) trifluoroborate (18.7 g, 74.6 mmol), Cs₂CO₃ (36.5 g, 112 mmol) in toluene (285 mL) and water (95 mL) was added PdCl₂(dppf) (1.37 g, 1.87 mmol)
- 15 under nitrogen protection. The mixture was stirred at 80 °C for 40 h. The resulting solution was quenched with water and extracted with EtOAc. The combined organic layers were dried over Na₂SO₄ and filtered. The filtrate was concentrated under reduced pressure and the residue was purified by column chromatography over silica gel (eluting with a 1%-40% gradient of EtOAc in PE) to afford **Int-2fe-1**. LC/MS: (M+Na)⁺= 471.4.

20 *Step F – Synthesis of Intermediate Int-2f-1*

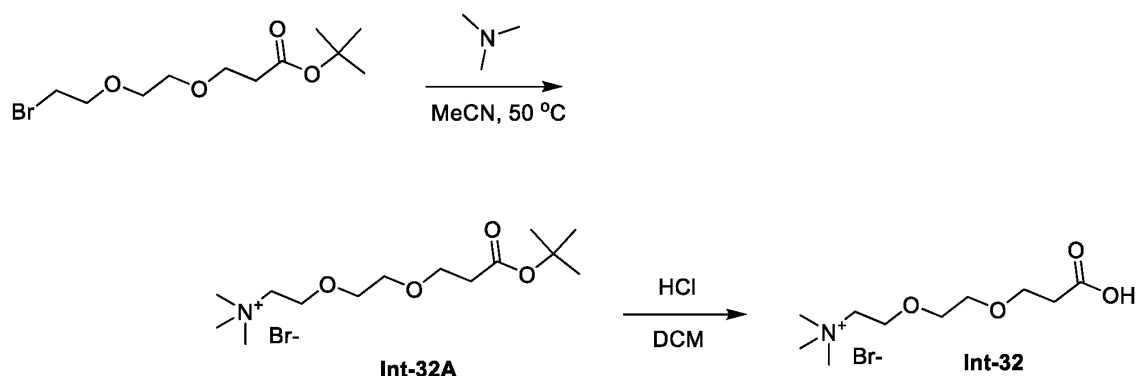
- To a solution of **Int-2fe1** (10.6 g, 22.4 mmol) in THF (100 mL) was added 1N TBAF in THF (44.9 mL, 44.9 mmol). This mixture was stirred at room temperature for 16 h then quenched with water and extracted with EtOAc. The combined organic layers were washed with brine, dried over anhydrous Na₂SO₄ and filtered. The filtrate was concentrated under
- 25 reduced pressure and the residue was purified by column chromatography over silica gel

(eluting with a 1%-70% gradient of EtOAc in PE) then by column chromatography over C18 (Column: 330 g; Mobile Phase A: water (10 mM NH₄HCO₃), Mobile Phase B: ACN; Flow rate: 80 mL/min; Gradient: 10% B to 10% B in 10 min, 20% B to 45% B in 10 min, 45% B to 70% B in 20 min Detector: UV 210 nm; Rt= 25 min) to provide **Int-2f-1**. LC/MS: (M+H)⁺= 305.1. ¹H NMR (300 MHz, CD₃OD) δ 7.27-7.16 (m, 4H), 5.85-5.75 (m, 1H), 5.00-4.85 (m, 2H), 3.79 (t, J = 7.0 Hz, 1H), 3.32-3.21 (m, 2H), 2.75 (t, J = 7.4 Hz, 2H), 2.98-1.72 (m, 4H), 1.42 (s, 9H).

EXAMPLE 2 Preparation of Ex-50 and Ex-52



The compound **Ex-50** is prepared in accordance with the scheme below from compound **Ex-01**, the preparation of which is described herein in Example 1, by reacting it under appropriate conditions with intermediate **Int 32**, prepared in accordance with the following Scheme:



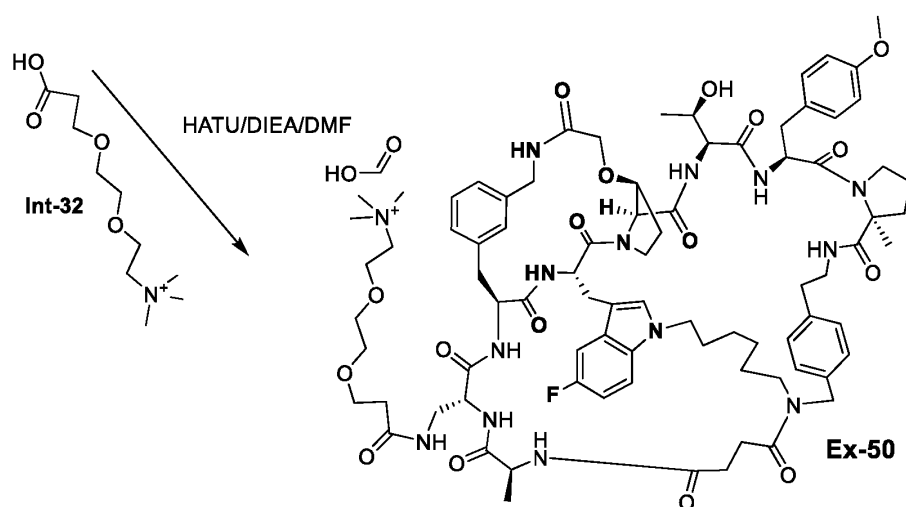
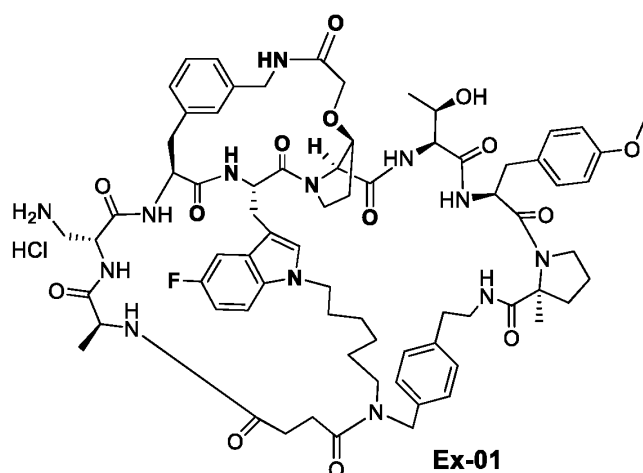
*Step A: Preparation of intermediate **Int-32A***

To a solution of tert-butyl 3-(2-(2-bromoethoxy)ethoxy)propanoate (5 g, 16.82 mmol) in acetonitrile (10 ml) was added trimethylamine (33% in ethanol, 11.46 ml, 50.5 mmol), and
 5 the resulting solution was heated at 50 °C overnight. The solution was concentrated to give 2-(2-(3-(tert-butoxy)-3-oxopropoxy)ethoxy)-N,N,N-trimethylethanaminium bromide (**Int-32A**).
 LC/MS: (M)⁺: 276.5.

*Step B: Preparation of intermediate **Int-32***

To a solution of 2-(2-(3-(tert-butoxy)-3-oxopropoxy)ethoxy)-N,N,N-trimethylethanaminium
 10 bromide (**Int-32A**) (5.99 g, 16.81 mmol) in DCM (20 ml) was added HCl (4N in dioxane) (21.01 ml, 84 mmol), and the resulting solution was stirred at rt overnight. The solution was concentrated to give 2-(2-(2-carboxyethoxy)ethoxy)-N,N,N-trimethylethanaminium bromide (**Int-32**). LC/MS: (M)⁺: 220.1.

Preparation of example compound Ex-50

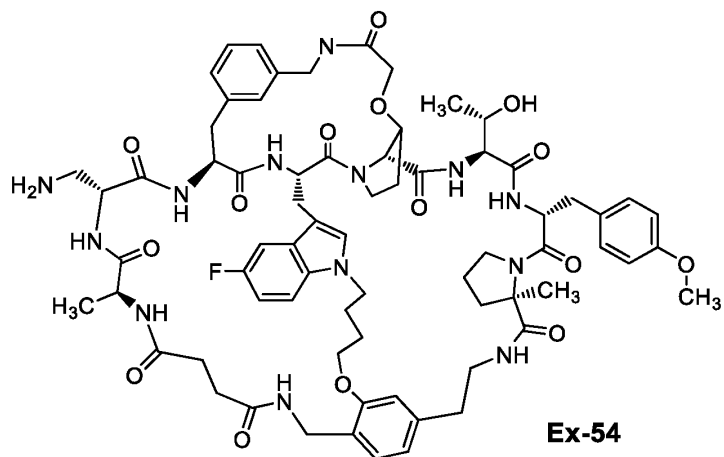
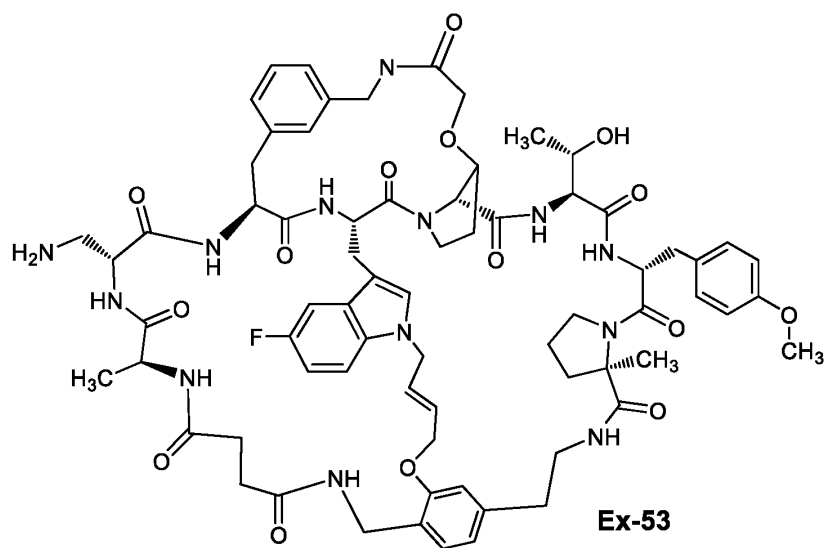


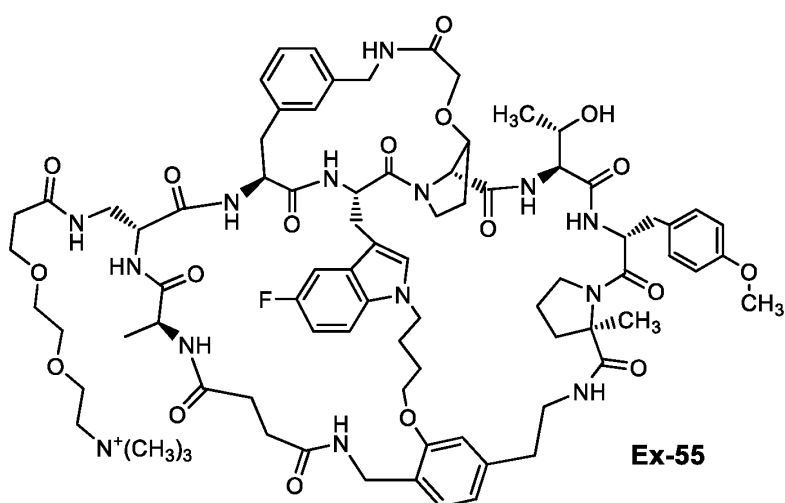
To a solution of **Ex-01** (crude) (17.4 mg, 0.012 mmol) and 2-(2-(2-carboxyethoxy)ethoxy)-N,N,N-trimethylethanaminium bromide (**Int-32**) (4.49 mg, 0.015 mmol) in DMF (2 ml) was added HATU (5.69 mg, 0.015 mmol) and DIEA (6.54 μ l, 0.037 mmol), and the resulting solution was stirred at rt for 50 min, then purified by reverse phase HPLC using acetonitrile(0.1%formic acid)/water(0.1%formic acid) as mobile phase to give **Ex-50**. LC/MS: M^+ = 1596.3.

Preparation of example compound Ex-52

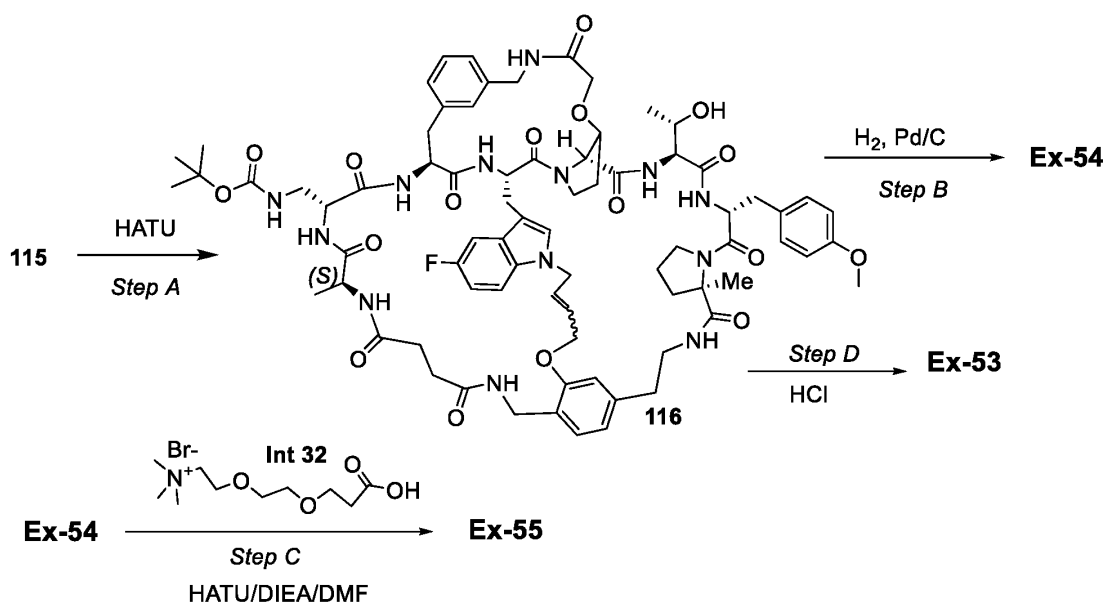
Compound **Ex-52** was prepared in an analogous manner to the preparation of compound **Ex-50** but using **Ex-51** instead of **Ex-01**. **Ex-52** was purified using reverse phase HPLC, in accordance with the methods described herein. LC/MS: M^+ = 1593.8.

EXAMPLE 3 Preparation of Ex-53, Ex-54 and Ex-55





Compounds **Ex-53**, **Ex-54** and **Ex-55** were prepared in a manner analogous to the compounds described above from intermediate **115** (preparation described below), according to the following schemes and synthesis description:



Step A – Synthesis of Intermediate **116**

To a solution of **115** (66.3 mg, 0.044 mmol) in DMF (1.5 ml), DCM (10 ml) and water (0.5 ml) at 0 °C was added DIPEA (0.030 ml, 0.173 mmol) followed by HATU (18.50 mg, 0.049 mmol) and the mixture was stirred for 30 min. The mixture was concentrated in vacuo and directly purified by column chromatography over C18 (30 g, eluting with acetonitrile+0.05%

TFA/water+0.05% TFA 90:10 to 40:60) to provide **116** as a mixture of E and Z isomers as well as **116** as pure fractions of E or Z isomer. LC/MS (major isomer) LC/MS: M^+ = 1481.19; LC/MS (minor isomer): LC/MS: M^+ = 1480.

Step B – *Synthesis of Compound Ex-54*

- 5 A solution of **116** (25.1 mg, 0.017 mmol) and Pd-C 10% (3.61 mg, 3.39 μ mol) in MeOH (10 ml) was hydrogenated at 1 atm for 1 h. The reaction was filtered over Celite and concentrated. The residue was treated with DCM/TFA 1:1 for 30 min then concentrated, treated with 4N HCl in dioxane (100 μ L) then concentrated to provide **Ex-54** in the form of the HCl salt. LC/MS: M^+ = 1383.44.

10 Step C – *Synthesis of Compound Ex-55*

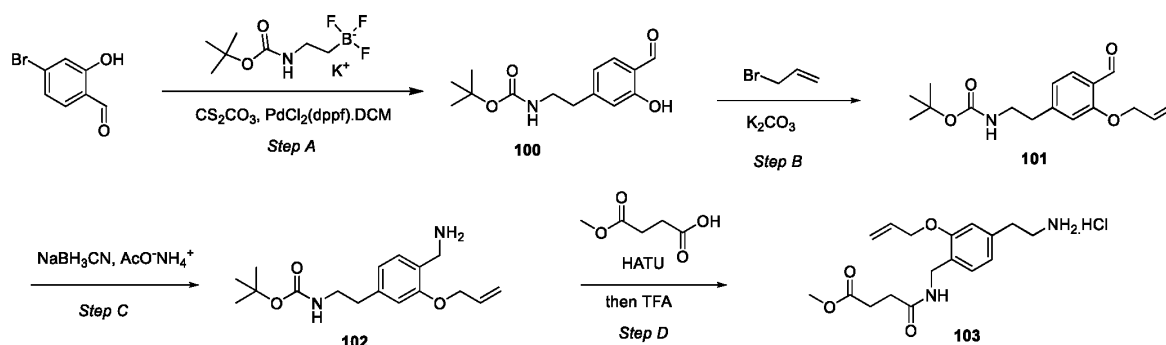
Example compound **Ex-55** was prepared in the form of the formate salt from **Ex-54** in a manner identical to that described in Example 2 for the synthesis of Compound **Ex-50**. LC/MS: M^+ = 1583.69.

Step D – *Synthesis of Compound Ex-53*

- 15 Example compound **Ex-53** was prepared in the form of the HCl salt from intermediate compound **116** in a manner identical to that described in Example 1 for the synthesis of compound **Ex-51**. LC/MS: M^+ = 1381.33.

Preparation of the intermediates needed to provide intermediate compound **115** from which compounds **Ex-53**, **Ex-54**, and **Ex-55** are ultimately prepared is described in the schemes and synthesis below beginning with the preparation of intermediate compound

20 **103**.



Step A – Synthesis of Intermediate **100**

A solution of 4-bromo-2-hydroxybenzaldehyde (3.00 g, 14.92 mmol), potassium tert-butyl N-[2-trifluoroboraniudyl]ethyl] carbamate (3.82 g, 15.22 mmol), cesium carbonate (17.02 g, 52.2 mmol) and 1,1'-bis(diphenylphosphino)ferrocene-palladium(ii)dichloride dichloromethane complex (0.611 g, 0.746 mmol) in degassed toluene (45 ml) and water (15 ml) was warmed to 75 °C and stirred overnight. The mixture was quenched at room temperature with half-saturated aqueous sodium bicarbonate and extracted with EtOAc. The combined organic fractions were washed with brine, dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with Hexanes/EtOAc 99:1 to 60:40) to give **100**. LC/MS: (M-55)⁺ = 210.25.

Step B – Synthesis of Intermediate **101**

To a solution of **100** (1.70 g, 6.41 mmol) and allyl bromide (0.832 ml, 9.61 mmol) in DMF (10 ml) at room temperature was added potassium carbonate (1.328 g, 9.61 mmol) and the mixture was warmed to 50 °C and stirred for 1 h. The mixture was quenched at room temperature with half-saturated aqueous sodium bicarbonate and extracted with EtOAc. The combined organic fractions were washed with brine, dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with Hexanes/EtOAc 99:1 to 70:30) to give **101**. LC/MS: (M-55)⁺ = 250.29.

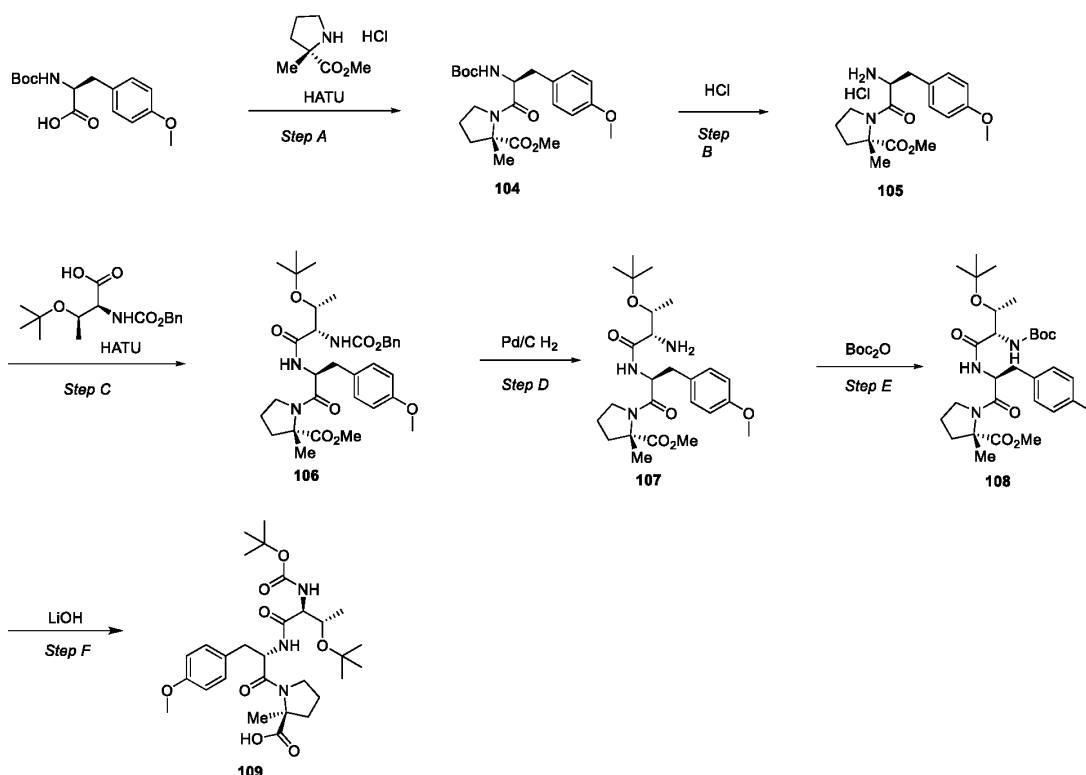
Step C – Synthesis of Intermediate **102**

To a solution of **101** prepared in the previous step (1.76 g, 5.76 mmol), 4 Å molecular sieves (2 g) and ammonium acetate (4.44 g, 57.6 mmol) in MeOH (100 ml) at room temperature was added sodium cyanoborohydride (0.380 g, 6.05 mmol) and the mixture was shaken overnight. The mixture was concentrated, quenched at room temperature with water and extracted with DCM. The combined organic fractions were dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with DCM/MeOH 99:1 to 30:70) to provide **102**. LC/MS: (2M+H)⁺ = 613.56.

Step D – Synthesis of Intermediate 103

To a solution of **102** (220 mg, 0.718 mmol) and succinic acid mono-methyl ester (114 mg, 0.862 mmol) in DMF (4 ml) at room temperature was added HATU (300 mg, 0.790 mmol) and DIPEA (0.314 ml, 1.795 mmol) and the mixture was stirred for 30 min. The mixture was quenched at room temperature with saturated aqueous sodium bicarbonate and extracted with EtOAc. The combined organic fractions were washed with brine, dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with Hexanes/EtOAc 99:1 to 30:70) to give an intermediate that was treated with 20% TFA in DCM for 1h. The reaction was concentrated then treated with 4N HCl (1.5 mL) and concentrated to provide **103**. LC/MS: (M+H)⁺ = 321.29.

Preparation of intermediate 109



Step A – Synthesis of Intermediate **104**

To a stirred solution of methyl (S)-2-methylpyrrolidine-2-carboxylate hydrochloride (7.00 g, 39 mmol) and (S)-2-((tert-butoxycarbonyl)amino)-3-(4-methoxyphenyl)propanoic acid (12.08 g, 40.9 mmol) in DMF (100 ml) at 0 °C was added DIPEA (17.01 ml, 97.0 mmol) followed by HATU (19.26 g, 50.7 mmol). The resulting mixture was allowed to warm to room temperature and stirred overnight. The reaction was quenched with a 10% aqueous LiCl solution and extracted with EtOAc. The organic extract was washed with 10% aqueous LiCl and dried over MgSO₄. The solvent was removed under reduced pressure and the residue purified by column chromatography over silica gel (eluting with Hexanes/EtOAc 80:20 to 40:60) to provide **104**.

Step B – Synthesis of Intermediate **105**

To a solution of **104** (16.4 g, 39.0 mmol) in EtOAc (100 ml) was added 4 N HCl in dioxane (48.8 ml, 195 mmol). The resulting mixture was stirred at room temperature for 18 h and was concentrated under reduced pressure to give **105** that was used in the next step without further purification.

5 *Step C – Synthesis of Intermediate 106*

To a solution of **105** (13.2 g, 37.0 mmol) and N-((benzyloxy)carbonyl)-O-(tert-butyl)-L-threonine (18.15 g, 37.0 mmol) in DMF at 0 °C was added DIPEA (16.15 ml, 92 mmol) followed by HATU (18.28 g, 48.1 mmol). The resulting mixture was allowed to warm to room temperature and stirred overnight. The reaction was quenched with 10% aqueous
10 LiCl solution and extracted with EtOAc. The organic extract was washed with 10% aqueous LiCl and dried over MgSO₄. The solvent was removed under reduced pressure and the residue purified by column chromatography over silica gel (eluting with Hexanes/EtOAc 80:20 to 40:60) to provide **106**.

Step D – Synthesis of Intermediate 107

15 To a solution of **106** (16.5 g, 27.0 mmol) in MeOH was added a slurry of 10% Pd/C and the mixture was hydrogenated at 20 psi for 4 h. The reaction mixture was filtered over Celite and concentrated under reduced pressure. The crude product was then re-dissolved in DCM and the solution was filtered through a 2 µm filter and concentrated to give **107**.

Step E – Synthesis of Intermediate 108

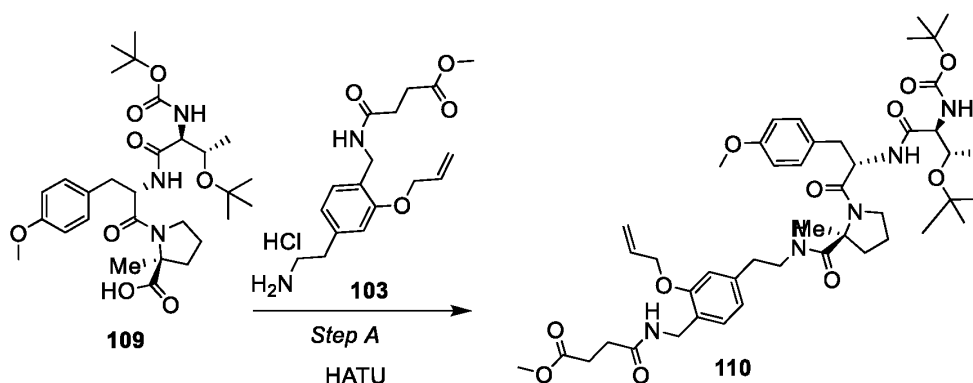
20 To a solution of **107** (3.2 g, 6.70 mmol) in DCM was added DIPEA (1.52 ml, 8.71 mmol) followed by di-tert-butyl dicarbonate (1.90 g, 8.71 mmol). The resulting mixture was stirred at room temperature for 2 h then concentrated under reduced pressure. The residue was purified by column chromatography over silica gel (eluting with Hexanes/EtOAc 100:0 to 40:60) to afford **108**.

25 *Step F – Synthesis of Intermediate 109*

A solution of **108** (1.43 g, 2.475 mmol) and 1 N aqueous LiOH (9.90 ml, 9.90 mmol) in THF (15 ml) and MeOH (15 ml) was warmed to 45 °C and stirred for 4 h then at 32 °C for 48 h. The reaction was concentrated, quenched at 0 °C with 0.5 M aqueous hydrochloric acid until pH ~2-3 and extracted with EtOAc. The combined organic fractions were dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with Hexanes/EtOAc-EtOH 99:1 to EtOAc-EtOH 3:1) to give **109**. LC/MS: (M+H)⁺ = 564.49.

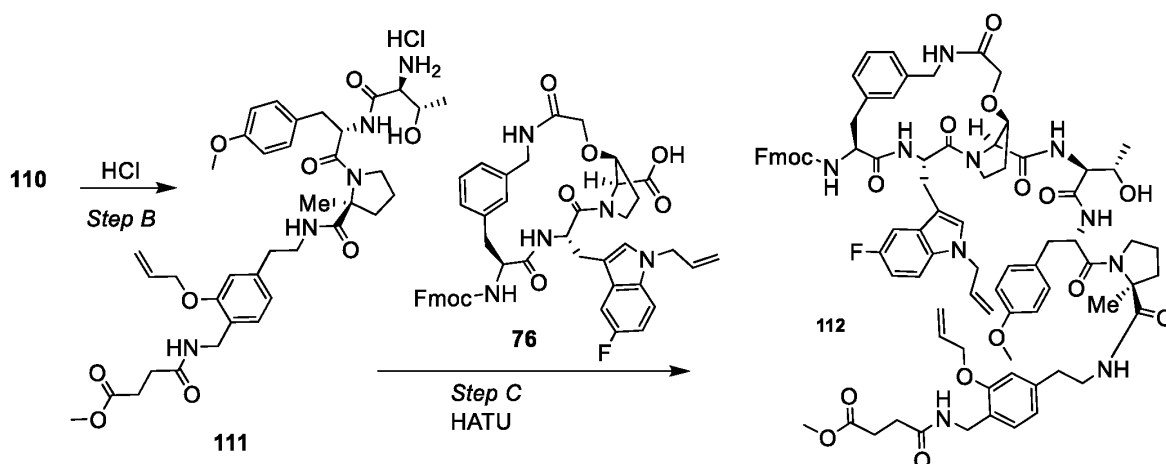
Preparation of Intermediate Compounds 110 to 115

Step A – Synthesis of Intermediate **110**



A solution of **109** (217 mg, 0.385 mmol), HATU (133 mg, 0.350 mmol) and DIPEA (0.245 ml, 1.400 mmol) in DMF (2.5 ml) was treated with **103** (217 mg, 0.385 mmol) at 0 °C and the mixture was allowed to warm to room temperature and stirred for 30 min. The mixture was quenched at 0 °C with saturated aqueous sodium bicarbonate and extracted with EtOAc. The combined organic fractions were washed with brine, dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with Hexanes/EtOAc-EtOH 3:1 99:1 to EtOAc-EtOH 3:1) to give **110**. LC/MS: (M+H)⁺ = 866.21.

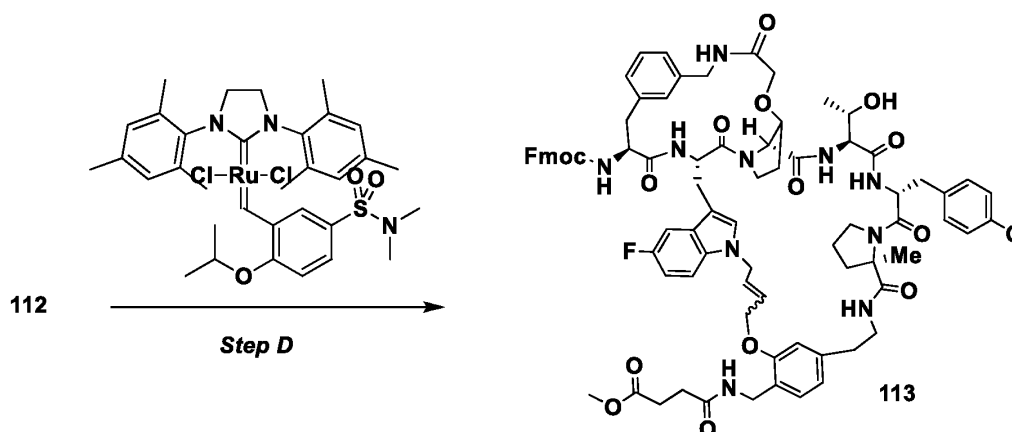
Steps B and C – Synthesis of Intermediate Compounds **111** and **112**



To a solution of **110** (249 mg, 0.288 mmol) in DCM (1 ml) at room temperature was added HCl 4 N in dioxane (0.359 ml, 1.438 mmol) and the mixture was stirred for 6 h then concentrated to provide **111**. LC/MS: (M+H)⁺ = 710.19.

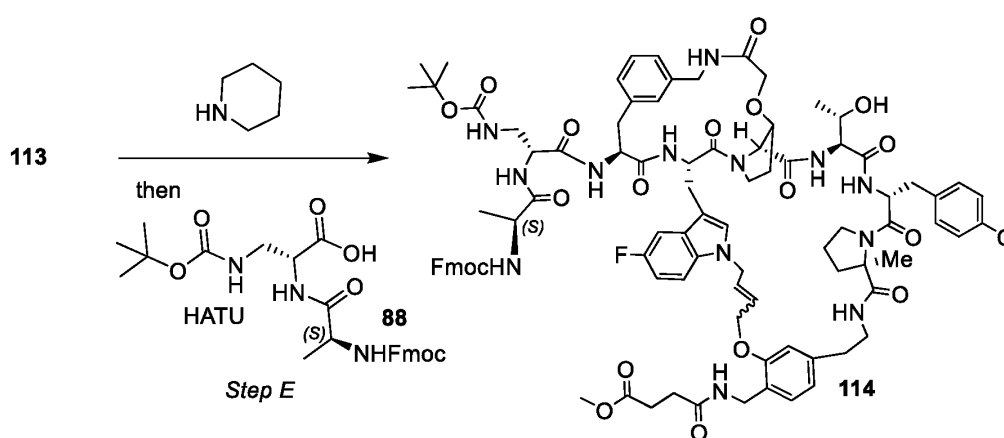
- 5 To a solution of **111** (219 mg, 0.293 mmol) and **76** (232 mg, 0.285 mmol) in DMF (3 ml) and water (0.15 ml) at 0 °C was added DIPEA (0.128 ml, 0.734 mmol) and HATU (123 mg, 0.323 mmol) and the mixture was stirred for 30 min. The mixture was quenched at 0 °C with brine and extracted with EtOAc. The combined organic fractions were dried over Na₂SO₄, filtered and concentrated in vacuo. The residue was purified by column
- 10 chromatography over silica gel (eluting with Hexanes/EtOAc-EtOH 3-1 99:1 to 30:70 then DCM/MeOH 99:1 to 70:30) to afford **112**. LC/MS: (M+H)⁺ = 1506.11.

Step D – Synthesis of Intermediate 113



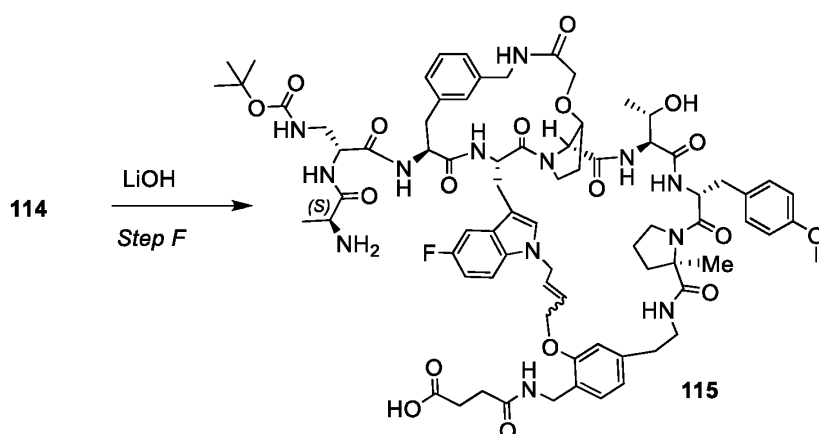
To a solution of **112** (173 mg, 0.115 mmol) in DCM (180 ml) and AcOH (15 ml) degassed with nitrogen for 30 min was added Zhan's Catalyst (59.0 mg, 0.080 mmol) and the mixture was warmed to 50 °C and stirred for 3 h. The mixture was filtered over Celite, washing with DCM then concentrated in vacuo. The residue was purified by column chromatography over silica gel (eluting with DCM/MeOH 99:1 to 80:20) to afford **113** as a mixture of E and Z isomers. LC/MS (major isomer): (M)⁺ = 1477.80; LC/MS (minor isomer): (M)⁺ = 1478.28.

Step E – Synthesis of Intermediate 114



To a solution of **113** (141 mg, 0.095 mmol) in acetonitrile (2 ml) was added piperidine (0.066 ml, 0.668 mmol) and the mixture was stirred for 45 min. The mixture was concentrated in vacuo, co-evaporated with acetonitrile trice to give a crude. To a slurry of this crude (119 mg, 0.095 mmol) and intermediate compound **88** (52.0 mg, 0.105 mmol) in DMF (2 ml) and water (0.1 ml) at 0 °C was added HATU (39.7 mg, 0.105 mmol) and DIPEA (0.037 ml, 0.209 mmol) and the mixture was stirred for 30 min. The mixture was purified by column chromatography over C18 (eluting with acetonitrile+0.05% TFA/water+0.05% TFA 90:10 to 30:70) to give **114** as a mixture of E and Z isomers. LC/MS major isomer: (M)⁺ = 1735.28; LC/MS (minor isomer): (M)⁺ = 1735.25.

Step F – Synthesis of intermediate compound 115



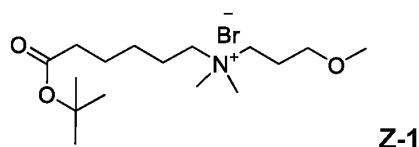
To a solution of **114** (134 mg, 0.077 mmol) in THF (1.5 ml) and MeOH (1.5 ml) at 0 °C was added 1 N aqueous LiOH (0.386 ml, 0.386 mmol) dropwise and the mixture was stirred for 2 h. The reaction was treated at 0 °C dropwise with 0.5 N HCl until pH~7, concentrated from organic solvents, then the slurry was dissolved with ~1 mL of DMF and directly purified by column chromatography over C18 (eluting with acetonitrile+0.05% TFA/water+0.05% TFA 90:10 to 50:50) to give **115** as a mixture of E and Z isomers. LC/MS major isomer: (M)⁺ = 1498.71; LC/MS (minor isomer): (M)⁺ = 1499.48.

As described above in the preparation of **Ex-50** from **Ex-01**, and **Ex-55** from **Ex-54** by reaction of the R² amide thereof in the respective starting compounds with an acidic substituent precursor, the following intermediate compounds may be employed in analogous reactions to provide useful compounds of the invention.

Synthesis of R¹/R² substituent precursors:

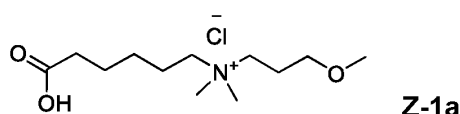
Preparation of 5-carboxy-N-(3-methoxypropyl)-N,N-dimethylpentan-1-aminium chloride (Intermediate Z-1a)

Step A: preparation of intermediate **Z-1**



To a stirred solution of *tert*-butyl 6-(dimethylamino)hexanoate (300 mg, 1.393 mmol) in acetonitrile (1 mL) was added 1-bromo-3-methoxypropane (853 mg, 5.57 mmol). The reaction mixture was stirred at 50 °C for 16 h. The resulting mixture was concentrated under reduced pressure to afford **Z-1**. LC/MS: (M-Br)⁺ = 288.4. ¹H NMR (300 MHz, CDCl₃):
5 δ 3.76-3.47 (m, 6H), 3.38 (d, *J* = 28.9 Hz, 9H), 2.25 (t, *J* = 7.2 Hz, 2H), 2.12-1.95 (m, 2H), 1.87-1.55 (m, 4H), 1.45 (s, 11H).

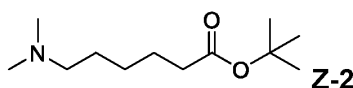
Step B: Synthesis of intermediate compound Z-1a



To a stirred solution of **Z-1** (460 mg, 1.249 mmol) in DCM (0.5 mL) was added 4 M HCl in
10 dioxane (2 mL) at room temperature. The reaction mixture was stirred at room temperature for 4 h and concentrated under reduced pressure. The residue was re-dissolved in DCM (5 mL) and concentrated under reduced pressure to afford intermediate compound **Z-1a**.
LC/MS: (M-Cl)⁺ = 232.3.

Preparation of Intermediate Z-2b

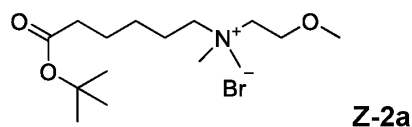
15 *Step A: preparation of intermediate Z-2*



To a stirred solution of *tert*-butyl 6-bromohexanoate (1.0 g, 3.98 mmol) in THF (10 mL) was added dimethylamine (2 M in THF) (7.96 mL, 15.93 mmol). The reaction mixture was stirred at room temperature for 16 h. The resulting mixture was concentrated under reduced
20 pressure and the residue was purified by silica gel column chromatography, eluting with a gradient 1% - 15% MeOH in DCM. The fractions containing the desired product were combined and concentrated to afford **Z-2**. LC/MS: (M+H)⁺ = 216.2. ¹H NMR (300 MHz,

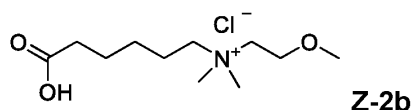
CDCl₃): δ 2.35-2.17 (m, J = 8.5, 6.5 Hz, 10H), 1.67-1.47 (m, 4H), 1.45 (s, 9H), 1.42 -1.23 (m, 2H).

Step B: preparation of intermediate **Z-2a**



To a stirred solution of **Z-2** (250 mg, 1.161 mmol) in ACN (1 mL) was added 1-bromo-2-methoxyethane (645 mg, 4.64 mmol). The reaction mixture was stirred at 50 °C for 16 h. The resulting mixture was concentrated under reduced pressure to afford **Z-2a**. LC/MS: (M-Br)⁺ = 274.3. ¹H NMR (300 MHz, CDCl₃): δ 4.02-3.80 (m, 4H), 3.70-3.54 (m, 2H), 3.42 (d, J = 13.0 Hz, 9H), 2.24 (t, J = 7.2 Hz, 2H), 1.85-1.75 (m, 2H), 1.72-1.55 (m, 2H), 1.44 (s, 11H).

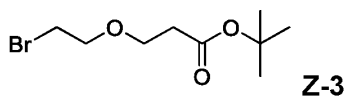
Step C: preparation of intermediate **Z-2b**



To a stirred solution of **Z-2a** (450 mg, 1.270 mmol) in DCM (0.5 mL) was added 4 M HCl in dioxane (2 mL) at room temperature. The reaction mixture was stirred at room temperature for 4 h and concentrated under reduced pressure. The residue was re-dissolved in DCM (5 mL) and concentrated under reduced pressure to afford **Z-2b**. LC/MS: (M-Cl)⁺ = 218.3.

Preparation of Intermediate Z-3b

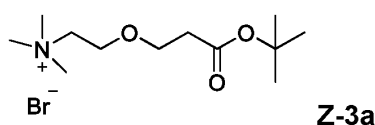
Step A: preparation of intermediate **Z-3**



To a solution of *tert*-butyl 3-(2-hydroxyethoxy)propanoate (500.0 mg, 2.63 mmol) in DCM (2 mL) were added CBr₄ (1395 mg, 4.21 mmol) and PPh₃ (965 mg, 3.68 mmol) at 0 °C. The mixture was stirred at room temperature for 2 h. The resulting mixture was concentrated under reduced pressure and the residue was purified by silica gel column chromatography,

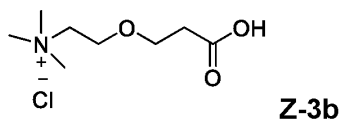
eluted with gradient 1% - 15% EA in PE. The fractions containing desired product were combined and concentrated to afford **Z-3**. ^1H NMR (400 MHz, CDCl_3): δ 3.78 (dt, J = 11.1, 6.3 Hz, 4H), 3.47 (t, J = 6.3 Hz, 2H), 2.53 (t, J = 6.4 Hz, 2H), 1.48 (s, 9H).

Step B: synthesis of intermediate Z-3a



To a stirred solution of tert-butyl 3-(2-bromoethoxy)propanoate **Z-3** (450 mg, 1.778 mmol) in ACN (2 mL) was added trimethylamine (955 mg, 5.33 mmol) (33%Wt, in EtOH). The reaction mixture was stirred at 50 °C for 16 h. The resulting mixture was concentrated under reduced pressure to afford **Z-3a**. LC/MS: (M-Br) $^+$ = 232.3. ^1H NMR (400 MHz, CDCl_3): δ 5.32 (s, 1H), 4.04-3.94 (m, 4H), 3.73 (t, J = 5.7 Hz, 2H), 3.50 (s, 10H), 2.50 (t, J = 5.7 Hz, 2H), 1.44 (s, 9H).

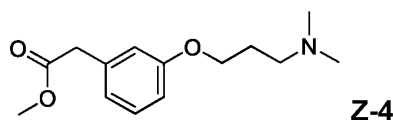
Step C: synthesis of intermediate Z-3b



To a solution of **Z-3a** (550 mg, 1.761 mmol) in DCM (0.6 mL) was added 4 M HCl in dioxane (2.5 mL) at room temperature. The mixture was stirred at room temperature for 4 h. The resulting mixture was concentrated under reduced pressure and the residue was re-dissolved in DCM (3 mL) and toluene (3 mL). The mixture was then concentrated under reduced pressure to afford **Z-3b**. LC/MS: (M-Cl) $^+$ = 176.2.

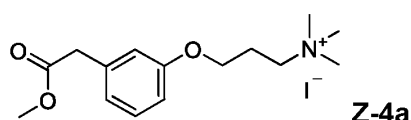
Preparation of Intermediate Z-4b

Step A: preparation of intermediate Z-4



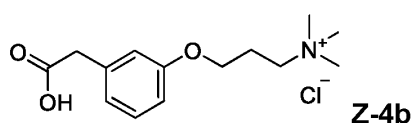
To a solution of DIAD (1.755 mL, 9.03 mmol) in THF (30 mL) was added Ph_3P (2.368 g, 9.03 mmol). The mixture was stirred at room temperature for 10 min, then methyl 2-(3-hydroxyphenyl)acetate (1.0 g, 6.02 mmol) and 3-(dimethylamino)propan-1-ol (0.931 g, 9.03 mmol) were added to the solution. The mixture was stirred at 50 °C for 1 h. The resulting solution was concentrated under reduced pressure and the residue was purified by silica gel column chromatography, eluting with a gradient 1% - 10% MeOH in DCM. The fractions containing the desired product were combined and concentrated to afford **Z-4**. LC/MS: $(\text{M}+\text{H})^+ = 252.2$. ^1H NMR (300 MHz, CDCl_3): δ 7.23-7.18 (m, 1H), 6.82 (td, $J = 8.7, 4.1$ Hz, 3H), 4.01 (t, $J = 6.4$ Hz, 2H), 3.69 (s, 3H), 3.59 (s, 2H), 2.46 (t, $J = 7.3$ Hz, 2H), 2.27 (s, 6H), 1.97 (dt, $J = 7.9, 6.5$ Hz, 2H).

Step B: Preparation of intermediate Z-4a



To a solution of **Z-4** (600 mg, 2.268 mmol) in ACN (12 mL) was added MeI (1.288 g, 9.07 mmol). The mixture was stirred at room temperature for 1 h. The resulting solution was concentrated under reduced pressure to afford **-Z-4a**. LC/MS: $(\text{M}-\text{I})^+ = 266.2$.

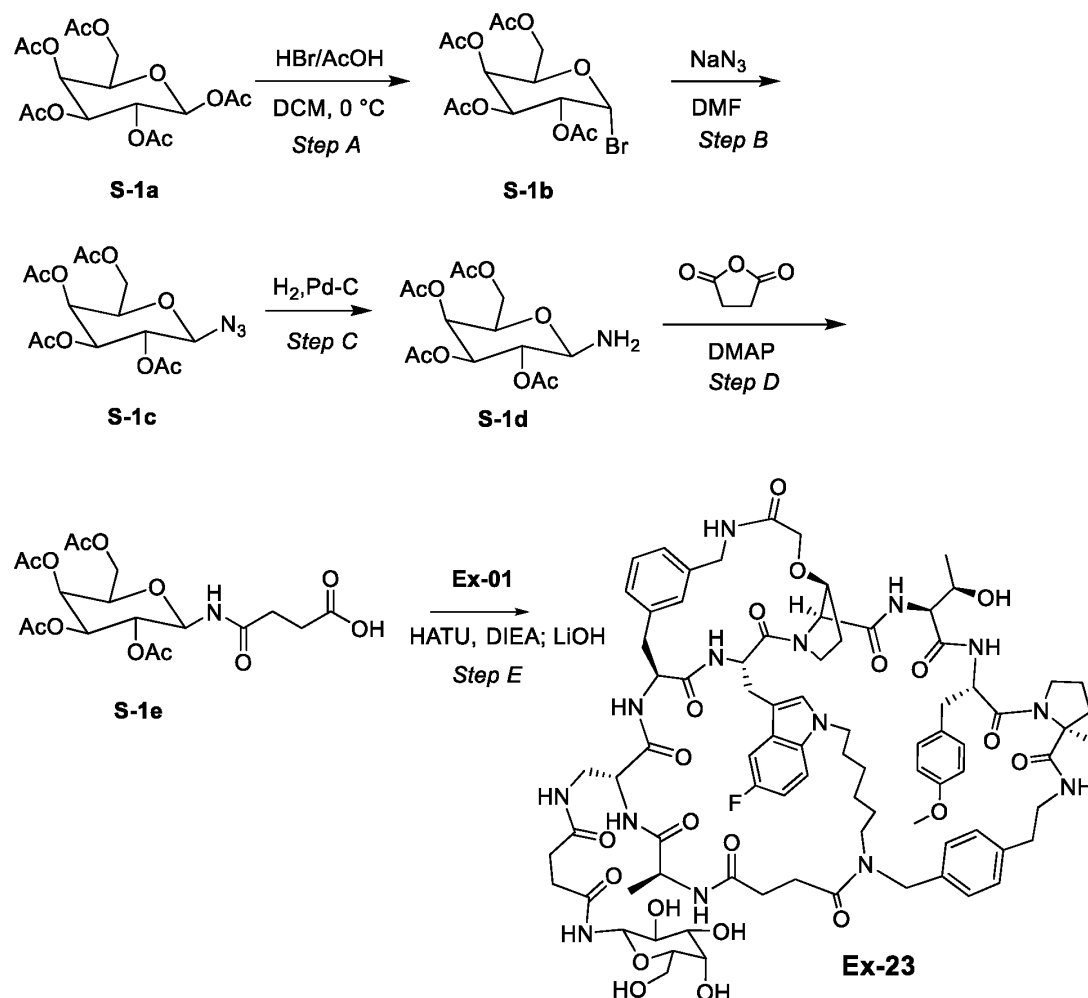
Step C: Preparation of intermediate Z-4b



To a solution of **Z-4a** (800 mg, 1.729 mmol) in THF (12 mL) was added 2 M LiOH (1.729 mL, 3.46 mmol). This mixture was stirred at room temperature for 2 h. The pH value of the solution was adjusted to 4 with HCl (1 M) and the solution was concentrated under reduced pressure. The crude product was purified by reverse phase chromatography over C18 (Mobile Phase A: water, Mobile Phase B: ACN; Flow rate: 60 mL/min; Gradient: 1% B to 25% B in 25 min; 25% B to 95% B in 15 min; 95% B to 95% B in 10 min) to afford **Z-4b**.

LC/MS: (M-Cl)⁺ = 252.2. ¹H NMR (300 MHz, CD₃OD): δ 7.21 (t, *J* = 7.9 Hz, 1H), 6.95-6.75 (m, 3H), 4.12 (t, *J* = 5.7 Hz, 2H), 3.63-3.50 (m, 4H), 3.18 (s, 9H), 2.35-2.20 (m, 2H).

EXAMPLE 4 Preparation of Ex-23



5

Step A: Preparation of intermediate **S-1b**

(2S,3R,4S,5S,6R)-6-(acetoxymethyl)tetrahydro-2H-pyran-2,3,4,5-tetraol/tetraacetate **S-1a** (5 g, 12.81 mmol) was added to a solvent of 48% HBr (7.25 mL, 64.0 mmol) in AcOH and DCM (40 mL) at 0°C and the mixture was stirred at 0°C for 1 hour. The reaction mixture
10 was poured into aqueous saturated sodium hydrogen carbonate cooled on ice and the mixture was extracted with DCM (3 x 60mL). The organic layer was washed with brine,

dried over Na₂SO₄, filtered and concentrated then the crude product was purified by flash chromatography over silica gel (eluting with 0~30% EtOAc/PE) to **S-1b**.

Step B: Preparation of intermediate S-1c

To a solution of **S-1b** (4.5 g, 10.94 mmol) in dry DMF (45 mL) was added sodium azide
5 (0.854 g, 13.13 mmol) and the reaction was stirred at 18 °C for 30 min. The reaction mixture was diluted with water (30 mL) and extracted with EtOAc (3 x 80 mL). The organic layer was dried over Na₂SO₄ and evaporated to dryness. The crude product was purified by flash chromatography over silica gel (eluting with 0~30% EtOAc/PE) to give **S-1c**.

Step C: Preparation of intermediate S-1d

10 To a solution of **S-1c** (3.15 g, 8.44 mmol) in EtOH (60 mL) were added 10% Pd-C (0.898 g, 0.844 mmol). The reaction vessel was purged from air and filled with H₂ under 50 psi. The reaction was stirred at 18 °C for 5 h. The reaction mixture was diluted with EtOAc, filtered through Celite, and concentrated, to give **S-1d** which was used for the next step.

Step D: Preparation of intermediate S-1e

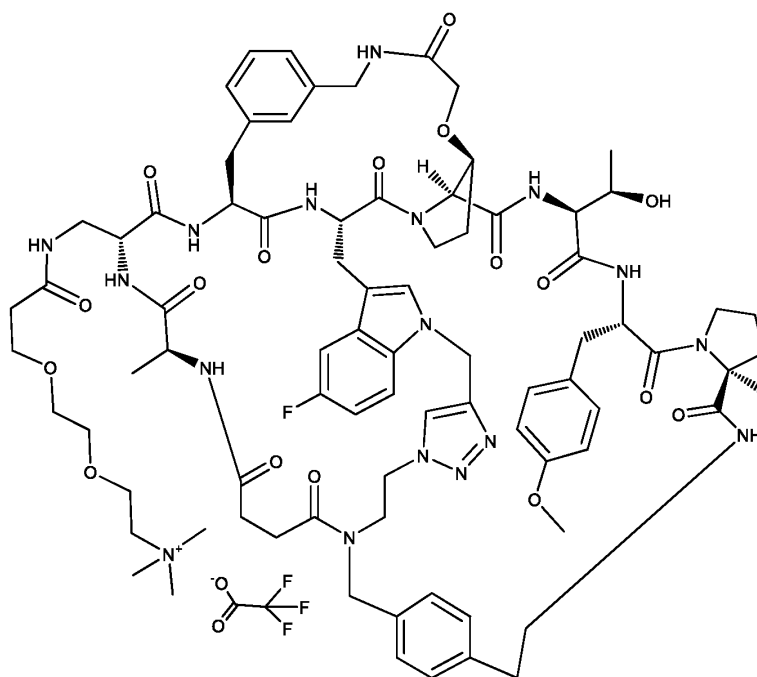
15 To a solution of **S-1d** (2.34 g, 6.74 mmol) in anhydrous THF (20 mL) was added dihydrofuran-2,5-dione (0.742 g, 7.41 mmol) and Et₃N (0.939 mL, 6.74 mmol). The reaction was stirred for 3 h until complete consumption of the starting material then evaporated. The resulting crude product was purified by flash chromatography over silica gel (eluting with 0~10% DCM/MeOH) to give **S-1e**. MS (ESI): m/z (M+H)⁺ 448.1. ¹H NMR
20 (400 MHz, CDCl₃) δ: 6.48 (d, J = 9.04 Hz, 1H), 5.43 (s, 1H), 5.24 (t, J = 8.93 Hz, 1H), 5.07-5.17 (m, 2H), 4.08-4.18 (m, 2H), 4.04 (q, J = 6.69 Hz, 1H), 2.72-2.84 (m, 1H), 2.58-2.69 (m, 2H), 2.43-2.53 (m, 2H), 2.15 (s, 3H), 2.06 (s, 3H), 2.04 (s, 4H), 2.00 (s, 3H).

Step E: Preparation of Ex-23

To a solution of **Ex-01** (300 mg, 0.215 mmol) and **S-1e** (115 mg, 0.258 mmol) in DMF (8 ml)
25 and water (0.4 ml) was added DIEA (0.150 ml, 0.860 mmol) and HATU (98 mg, 0.258

5

EXAMPLE 5 Preparation of Ex-14

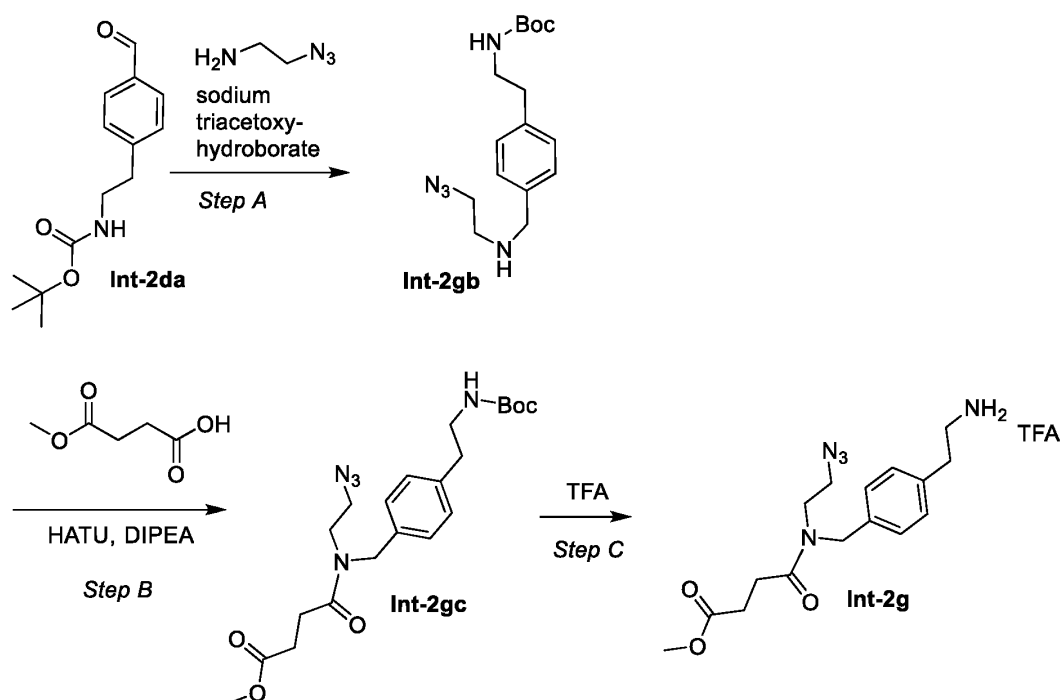


Ex-14

10

Preparation of intermediate Int-2g

Intermediate **Int-2g**, useful as a “linker” in the preparation of compounds of the invention, was prepared in accordance with the following scheme:



Step A – Synthesis of **Int-2gb**

To a solution of **Int-2da** (0.5 g, 2 mmol) and 2-azidoethanamine, HCl (0.246 g, 2mmol) in THF (16 ml) at room temperature in a water bath was added sodium triacetoxyhydroborate (1.06 g, 5 mmol) portion wise and the mixture was stirred for 2h. The reaction was slowly quenched with aqueous saturated NaHCO_3 solution, then extracted with DCM and washed with brine. The combined organic layers were dried over MgSO_4 and concentrated to give **Int-2gb**. LC/MS: $(\text{M}+1)^+ = 320.3$.

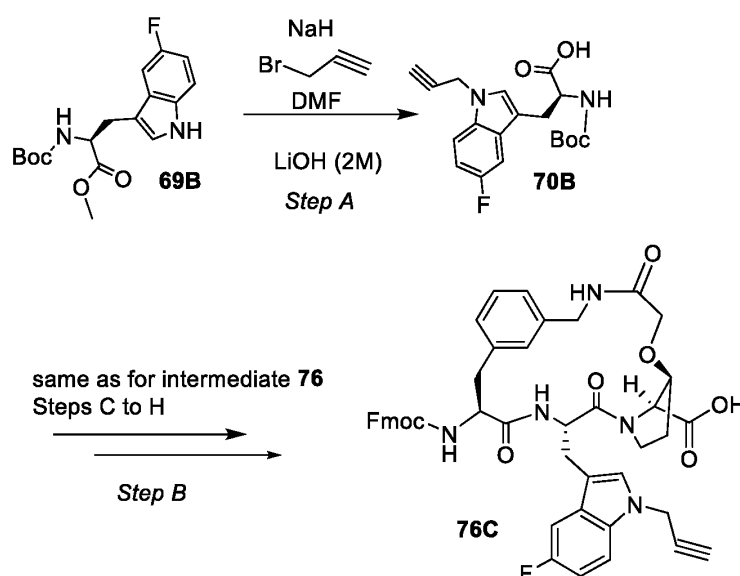
Step B – Synthesis of **Int-2gc**

To a solution of **Int-2gb** (0.64 g, 2 mmol) and mono-methyl succinate (0.3 g, 2.3 mmol) in DMF (4ml) and DCM (8 ml) was added HATU (0.914 g, 2.4 mmol) and DIPEA (0.7 ml, 4.01 mmol) at -15°C . The resulting solution was stirred at -15°C for 2 hours, then quenched with water and concentrated. The residue was purified by reverse-phase chromatography over C18 (eluting with acetonitrile/water + 0.1% TFA) to give **Int-2gc**. LC/MS: $(\text{M}+1)^+ = 434.3$.

Step C – Synthesis of Int-2g

To a solution of **Int-2gc** (0.52 g, 1.2 mmol) in DCM (9 mL) was added TFA (3 mL, 38.9 mmol) at room temperature. The reaction mixture was stirred at room temperature for 2 h. The mixture was concentrated under reduced pressure to afford **Int-2g**. LC/MS: (M+1)⁺=

5 334.3.

Preparation of intermediate compounds 70B and 76C*Step A – Synthesis of intermediate 70B*

To a solution of **69B** (1.5 g, 4.46 mmol) in DMF (17.8 ml) at 0 °C was added 95% NaH

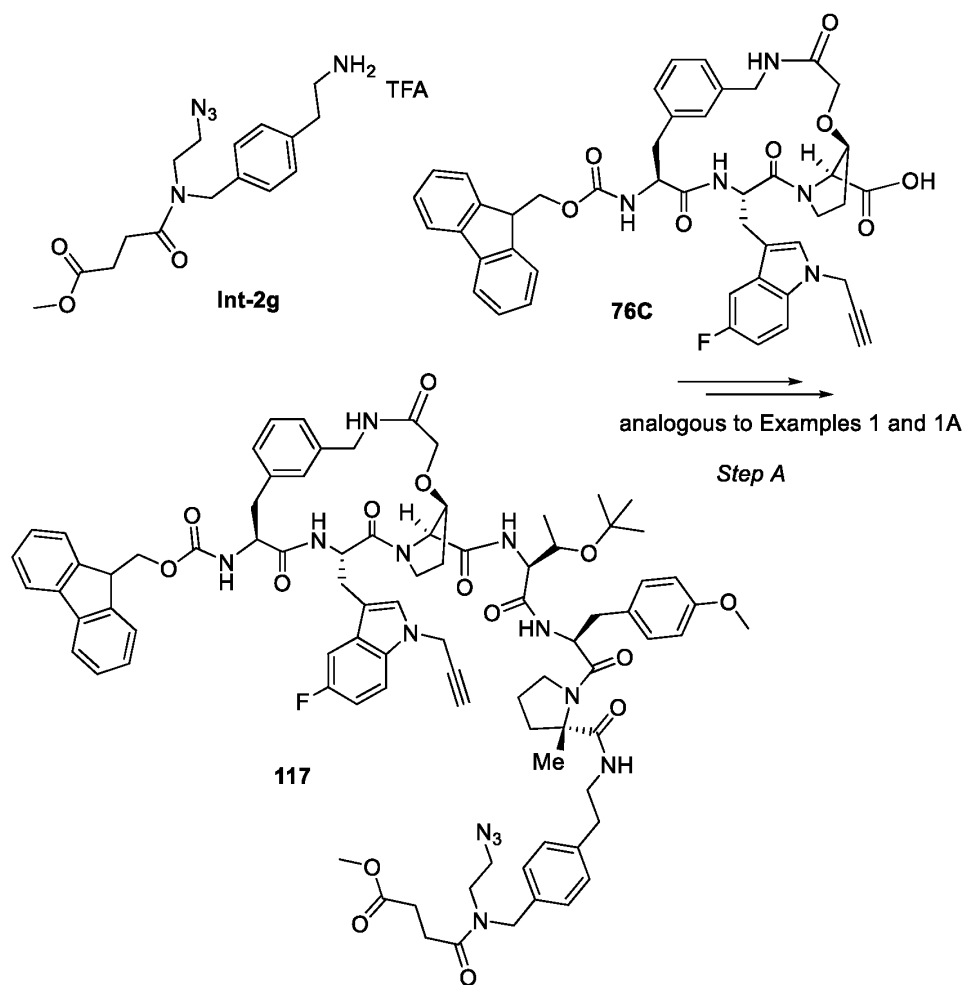
10 (0.141 g, 5.56 mmol), and the resulting solution was stirred at 0 °C for 20 min followed by addition of 3-bromoprop-1-yne (80% in toluene) (0.596 ml, 5.35 mmol) dropwise. To the resulting solution was added aqueous lithium hydroxide (2M) (3345 µl, 6.69 mmol) dropwise. The reaction was stirred at room temperature for 2h, filtered and purified by reverse phase HPLC (eluting with acetonitrile/water + 0.1% TFA) to give **70B**. LC/MS:

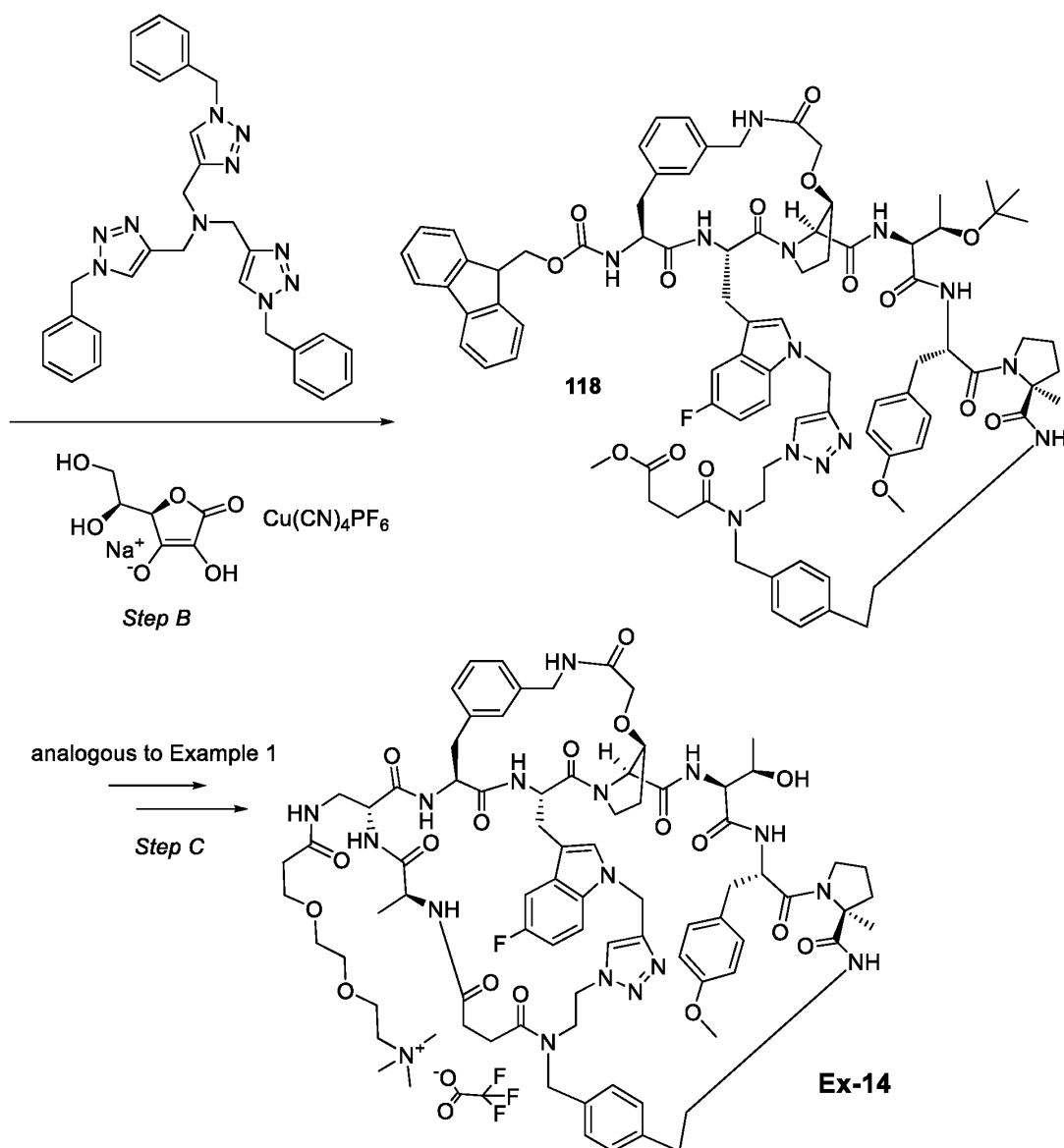
15 (M+1)⁺: 361.0, (M+Na)⁺: 383.0.

Step B – Synthesis of intermediate 76C

Conversion of **70B** to intermediate **76C** proceeded according to procedures analogous to those described in the preparation of intermediate **76** Steps C to H. LC/MS: $[M+1]^+ = 812.16$.

Assembly into Example Ex-14:





Step A – Synthesis of intermediate **117**

Intermediate **117** was prepared from intermediates **Int-2g** and **76C** according to procedures analogous to those described in Example 1 and 1A. More specifically **Int-2g** was

5 functionalized following reagents and procedures for the preparation of intermediate **77B**

Steps A to B, further elaborated following procedures for the preparation of intermediate **86**

Steps G to J, then finally coupled with intermediate **76C** following procedures for the

preparation of Example **1A** Steps B to C, to provide **117**. LC/MS: (M+1)⁺: 1573.36.

Step B – Synthesis of intermediate 118

Tetrakis(acetonitrile)copper(I) hexafluorophosphate (51.6 mg, 0.138 mmol), tris[(1-benzyl-1h-1,2,3-triazol-4-yl)methyl]amine (73.4 mg, 0.138 mmol) and sodium ascorbate (137 mg, 0.692 mmol) in BuOH (185.00 mL)/water (93 mL) were bubbled with nitrogen and then
5 heated at 50 °C. Intermediate **117** (435.3 mg, 0.277 mmol) was added into the reaction as a solid. After 1 h, the reaction was treated with aqueous pH 4 buffer and extracted with EtOAc. The combined organic layers were evaporated, and the residue was purified by reverse phase chromatography (eluting with a gradient of acetonitrile/water +0.1% formic acid) to provide intermediate **118**. LC/MS: (M+1)+: 1573.2.

10 *Step C – Synthesis of Ex-14*

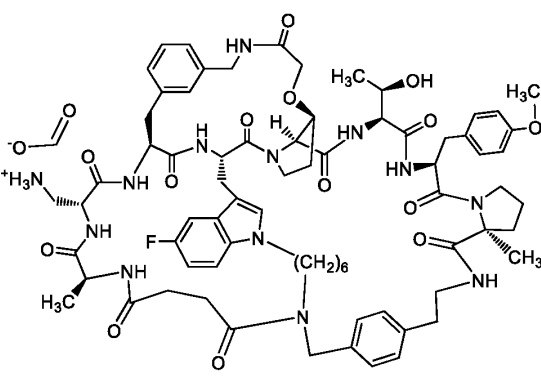
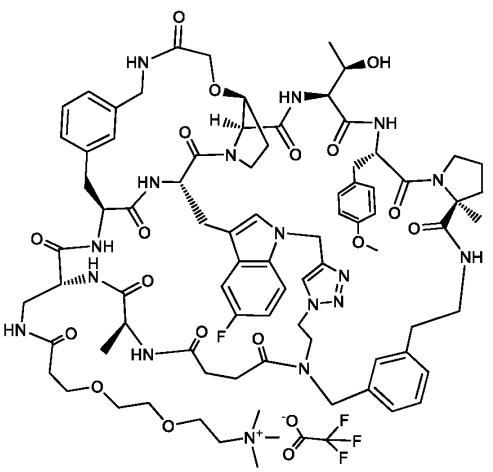
Synthesis of Example **Ex-14** proceeded from intermediate **118** according to procedures analogous to those described in Example 1, including the use of alternate spacers for the assembly. LC/MS: [M+1]+ = 1621.01.

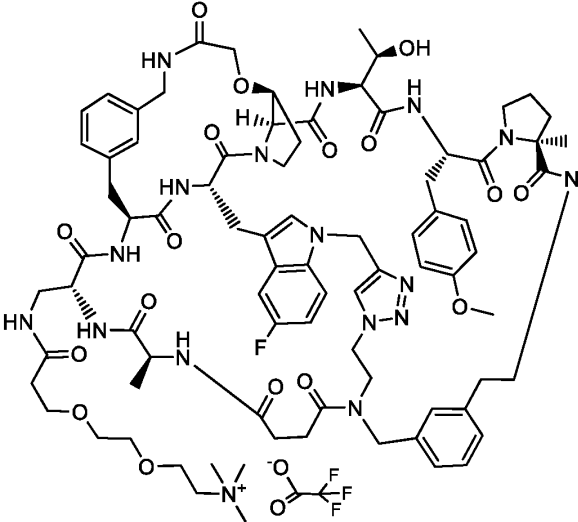
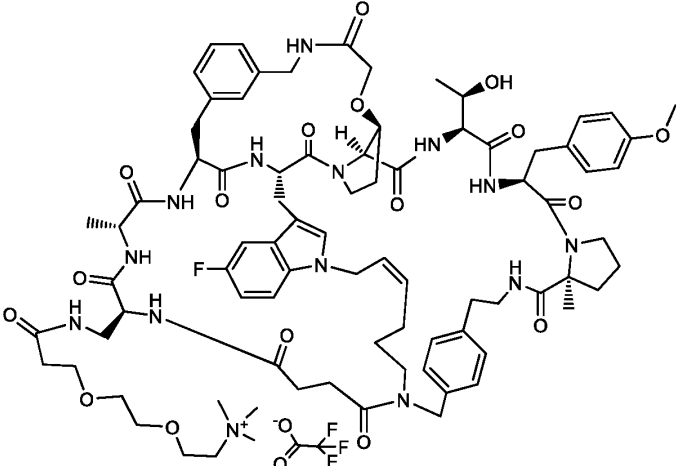
Using the synthetic schemes described above, and as will be appreciated, in some
15 instances with appropriate substitution of certain intermediates including the use of alternate spacers apparent to those skilled in the art, the preparation of which may be described above, the following compounds of the invention, listed in Table 2 below, were prepared. Additionally, alternate salt forms of compounds of the instant invention may also be described in this application:

20

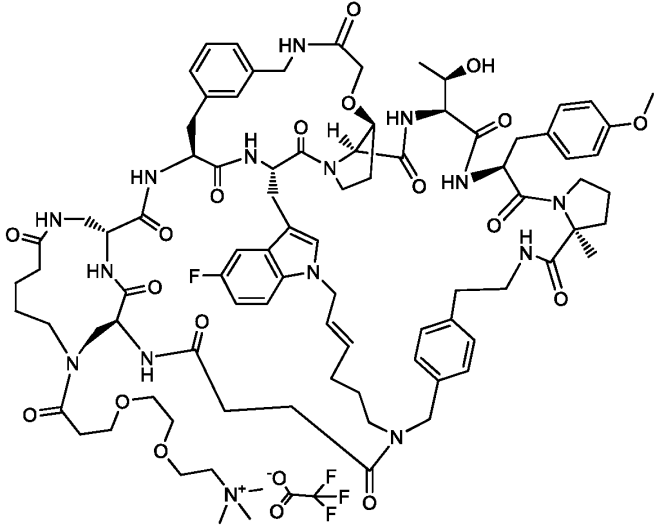
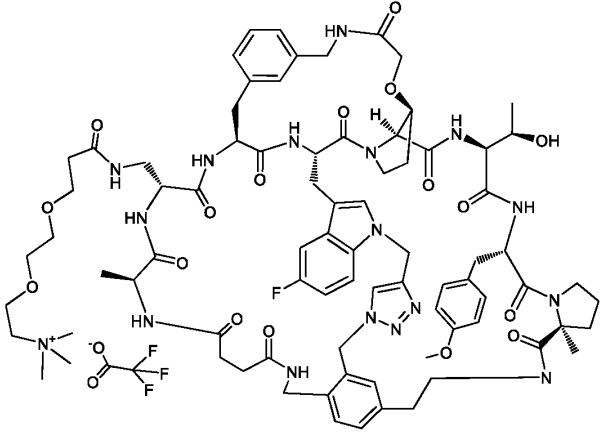
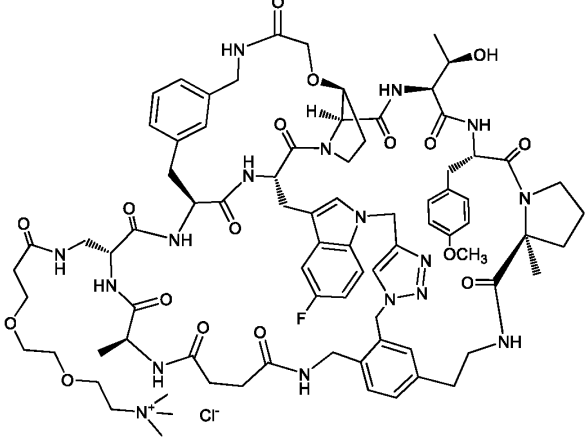
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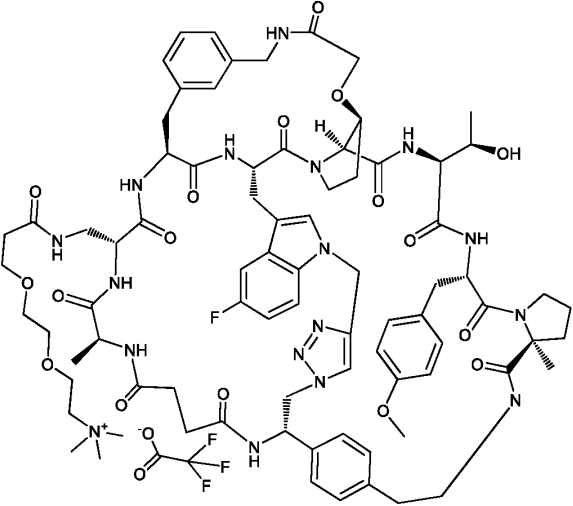
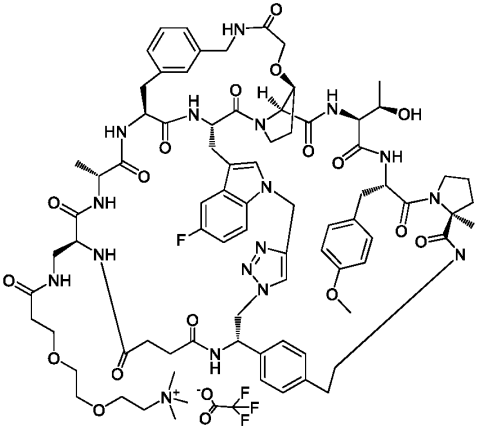
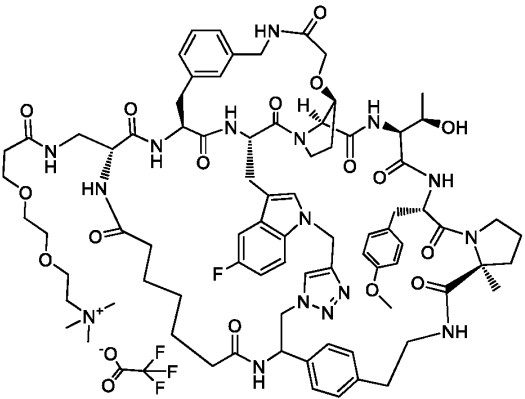
Table 2

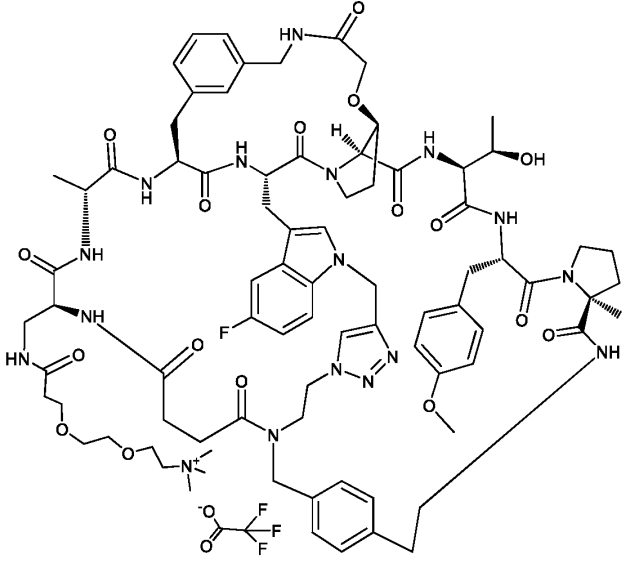
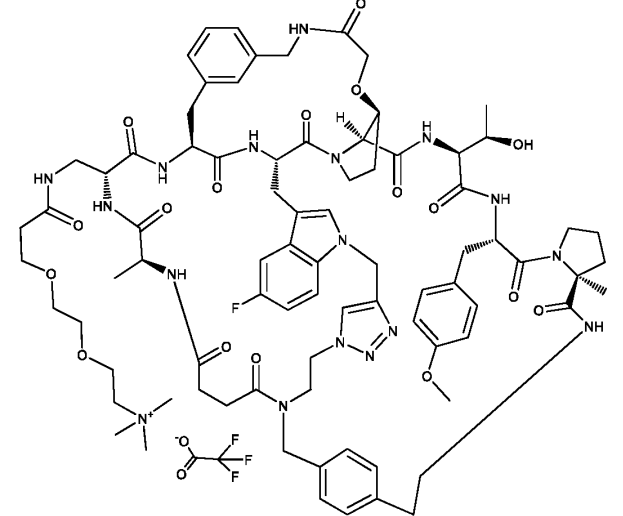
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-01 ACOH salt		1394.4
Ex-02 TFA salt		1622.12

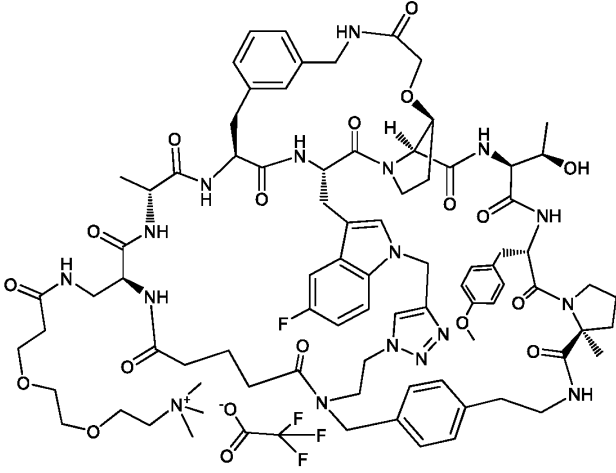
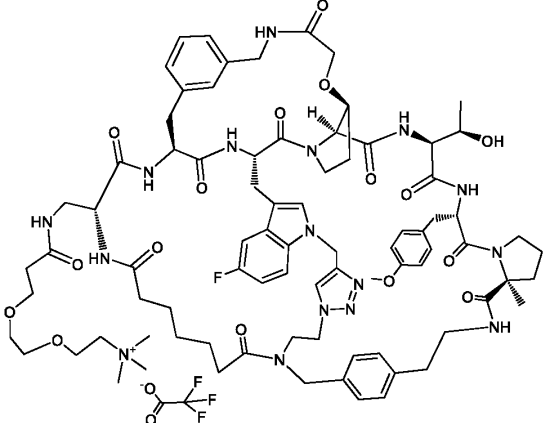
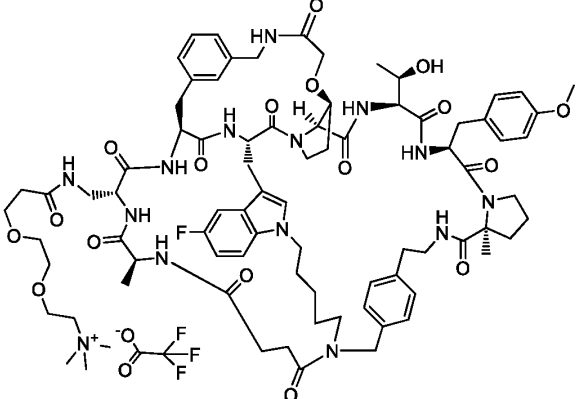
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-03 TFA salt	 The chemical structure of Ex-03 TFA salt is a complex macrocyclic molecule. It features a central benzimidazole core substituted with a 4-fluorophenyl group and a 4-methoxyphenyl group. The macrocycle is formed by several amide and ether linkages, including a 4-phenylbutyramide, a 2-hydroxy-3-methylbutyramide, a 2-methyl-2-oxazolidinone, and a 2-oxo-1,3-dioxane. A side chain contains a 4-phenylbutyramide and a 2-methyl-2-oxazolidinone. The molecule is shown as a TFA salt, with a trifluoroacetate anion (CF₃COO⁻) and a quaternary ammonium cation (N⁺Me₃) present.	1622.06
Ex-04 TFA salt	 The chemical structure of Ex-04 TFA salt is a complex macrocyclic molecule. It features a central benzimidazole core substituted with a 4-fluorophenyl group and a 4-methoxyphenyl group. The macrocycle is formed by several amide and ether linkages, including a 4-phenylbutyramide, a 2-hydroxy-3-methylbutyramide, a 2-methyl-2-oxazolidinone, and a 2-oxo-1,3-dioxane. A side chain contains a 4-phenylbutyramide and a 2-methyl-2-oxazolidinone. The molecule is shown as a TFA salt, with a trifluoroacetate anion (CF₃COO⁻) and a quaternary ammonium cation (N⁺Me₃) present.	1595.04

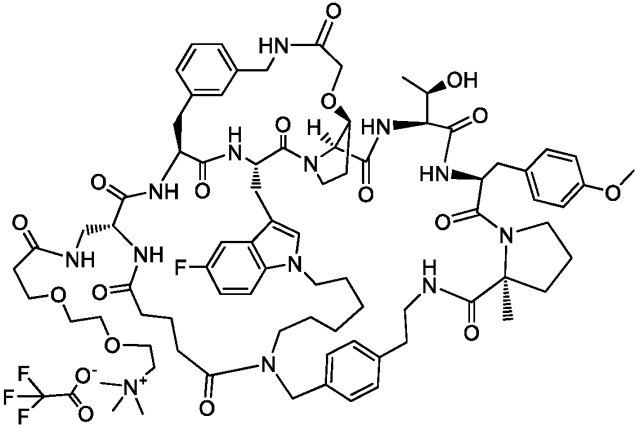
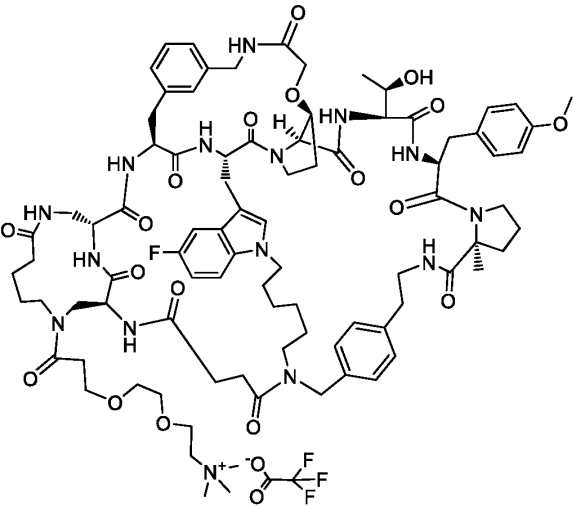
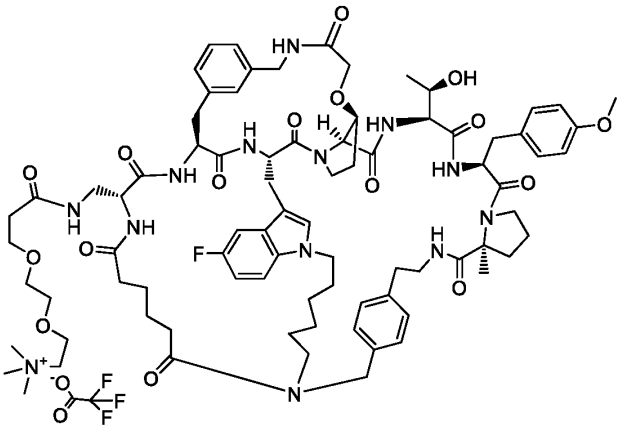
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-05 TFA salt		1594.54
Ex-06 TFA salt		1549.44

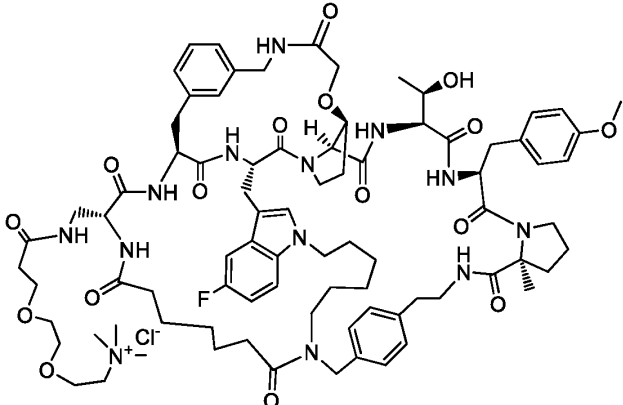
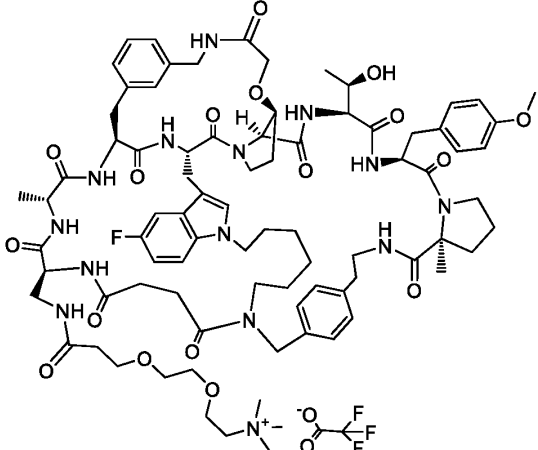
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-07 TFA salt	 The chemical structure of Ex-07 TFA salt is a large, complex molecule. It features a central indole ring system substituted with a fluorine atom and a long chain containing a double bond and a phenyl group. This central part is connected to various side chains, including a 4-methoxyphenyl group, a 4-methoxyphenyl group attached to a pyrrolidine ring, and a 4-methoxyphenyl group attached to a chiral center with a hydroxyl group. The molecule also contains a long chain with an ether linkage and a quaternary ammonium cation (N⁺) with a trifluoromethyl group (CF₃). The trifluoromethyl group is shown as a carbon atom bonded to three fluorine atoms and a nitrogen atom with a positive charge.	1692.66
Ex-08 TFA salt	 The chemical structure of Ex-08 TFA salt is a large, complex molecule. It features a central indole ring system substituted with a fluorine atom and a long chain containing a double bond and a phenyl group. This central part is connected to various side chains, including a 4-methoxyphenyl group, a 4-methoxyphenyl group attached to a pyrrolidine ring, and a 4-methoxyphenyl group attached to a chiral center with a hydroxyl group. The molecule also contains a long chain with an ether linkage and a quaternary ammonium cation (N⁺) with a trifluoromethyl group (CF₃). The trifluoromethyl group is shown as a carbon atom bonded to three fluorine atoms and a nitrogen atom with a positive charge.	1608.46
Ex-09 Cl salt	 The chemical structure of Ex-09 Cl salt is a large, complex molecule. It features a central indole ring system substituted with a fluorine atom and a long chain containing a double bond and a phenyl group. This central part is connected to various side chains, including a 4-methoxyphenyl group, a 4-methoxyphenyl group attached to a pyrrolidine ring, and a 4-methoxyphenyl group attached to a chiral center with a hydroxyl group. The molecule also contains a long chain with an ether linkage and a quaternary ammonium cation (N⁺) with a trifluoromethyl group (CF₃). The trifluoromethyl group is shown as a carbon atom bonded to three fluorine atoms and a nitrogen atom with a positive charge. A chloride ion (Cl⁻) is shown as a separate species.	1608.30

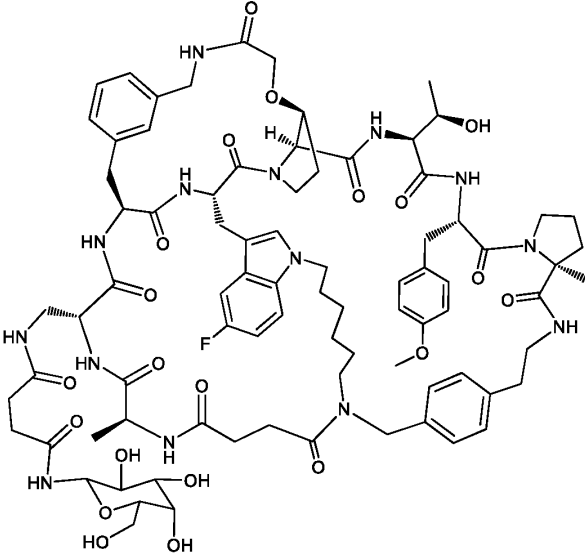
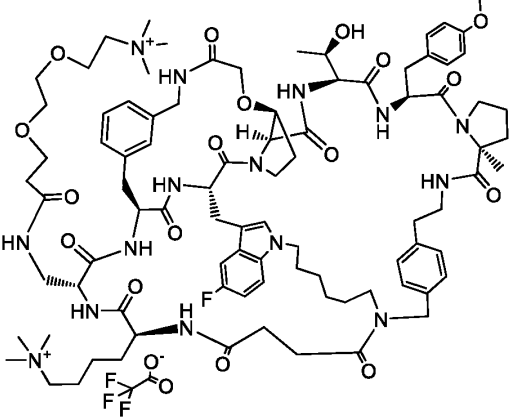
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-10 TFA salt	 The chemical structure of Ex-10 TFA salt is a large, complex molecule. It features a central indole ring system substituted with a fluorine atom and a triazole ring. The molecule is linked via various amide and ether bonds to several side chains, including a benzyl group, a 4-methoxyphenyl group, and a 4-fluorophenyl group. A trifluoromethyl group is also present. The molecule is shown as a cation with a trifluoromethyl sulfonate counterion.	1608.40
Ex-11 TFA salt	 The chemical structure of Ex-11 TFA salt is similar to Ex-10, but with a different arrangement of the side chains. It features a central indole ring system substituted with a fluorine atom and a triazole ring. The molecule is linked via various amide and ether bonds to several side chains, including a benzyl group, a 4-methoxyphenyl group, and a 4-fluorophenyl group. A trifluoromethyl group is also present. The molecule is shown as a cation with a trifluoromethyl sulfonate counterion.	1608.84
Ex-12 TFA salt	 The chemical structure of Ex-12 TFA salt is similar to Ex-10 and Ex-11, but with a different arrangement of the side chains. It features a central indole ring system substituted with a fluorine atom and a triazole ring. The molecule is linked via various amide and ether bonds to several side chains, including a benzyl group, a 4-methoxyphenyl group, and a 4-fluorophenyl group. A trifluoromethyl group is also present. The molecule is shown as a cation with a trifluoromethyl sulfonate counterion.	1579.54

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-13 TFA salt	 The chemical structure of Ex-13 TFA salt is a complex macrocyclic molecule. It features a central benzimidazole core. One side of the macrocycle contains a 4-methoxyphenyl group and a 4-fluorophenyl group. The other side includes a 4-methoxyphenyl group, a 4-fluorophenyl group, and a 4-methoxyphenyl group. The macrocycle is linked by amide bonds and includes a 4-methoxyphenyl group. The structure is shown as a TFA salt, with a trifluoroacetate anion (CF₃COO⁻) and a trimethylammonium cation (N⁺(CH₃)₃) present.	1621.74
Ex-14 TFA salt	 The chemical structure of Ex-14 TFA salt is a complex macrocyclic molecule, similar to Ex-13. It features a central benzimidazole core. One side of the macrocycle contains a 4-methoxyphenyl group and a 4-fluorophenyl group. The other side includes a 4-methoxyphenyl group, a 4-fluorophenyl group, and a 4-methoxyphenyl group. The macrocycle is linked by amide bonds and includes a 4-methoxyphenyl group. The structure is shown as a TFA salt, with a trifluoroacetate anion (CF₃COO⁻) and a trimethylammonium cation (N⁺(CH₃)₃) present.	1622.28

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-15 TFA salt	 The chemical structure of Ex-15 TFA salt is a complex macrocyclic molecule. It features a central benzimidazole core. Attached to this core are several side chains, including a 4-phenylbutyl group, a 2-hydroxypropyl group, and a 4-methoxyphenyl group. The molecule also contains a 1,3-dioxolane ring and a 1,3,4-oxadiazole ring. A trifluoroacetate (TFA) counterion is present, shown as a trifluoroacetate anion (CF₃COO⁻) and a quaternary ammonium cation (N⁺Me₃).	1636.54
Ex-16 TFA salt	 The chemical structure of Ex-16 TFA salt is a complex macrocyclic molecule, similar to Ex-15. It features a central benzimidazole core. Attached to this core are several side chains, including a 4-phenylbutyl group, a 2-hydroxypropyl group, and a 4-methoxyphenyl group. The molecule also contains a 1,3-dioxolane ring and a 1,3,4-oxadiazole ring. A trifluoroacetate (TFA) counterion is present, shown as a trifluoroacetate anion (CF₃COO⁻) and a quaternary ammonium cation (N⁺Me₃).	1592.52
Ex-17 TFA salt	 The chemical structure of Ex-17 TFA salt is a complex macrocyclic molecule, similar to Ex-15 and Ex-16. It features a central benzimidazole core. Attached to this core are several side chains, including a 4-phenylbutyl group, a 2-hydroxypropyl group, and a 4-methoxyphenyl group. The molecule also contains a 1,3-dioxolane ring and a 1,3,4-oxadiazole ring. A trifluoroacetate (TFA) counterion is present, shown as a trifluoroacetate anion (CF₃COO⁻) and a quaternary ammonium cation (N⁺Me₃).	1582.68

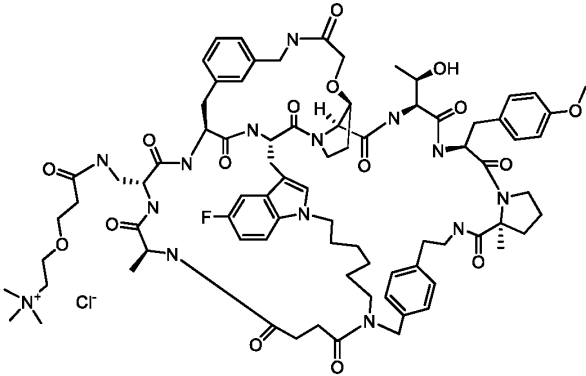
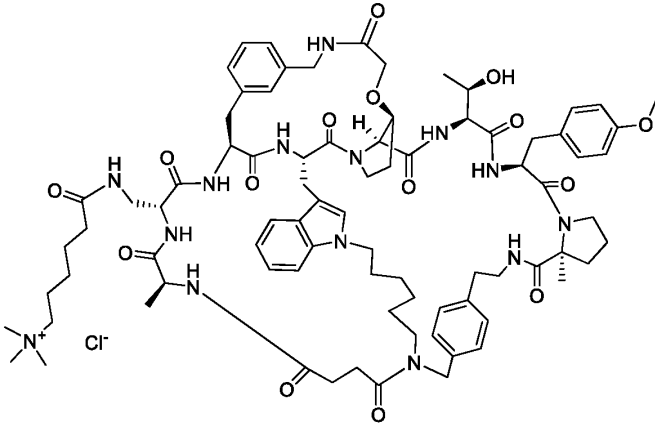
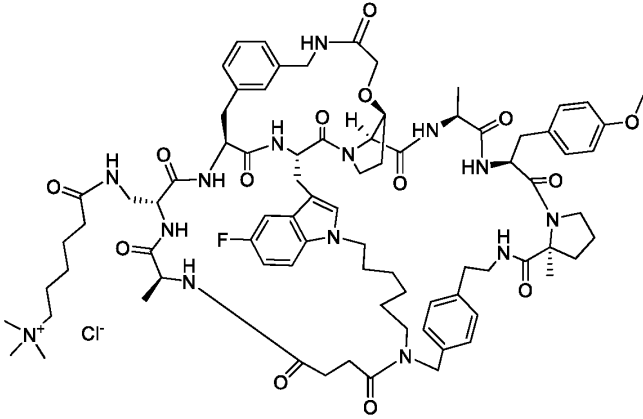
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-18 TFA salt	 The chemical structure of Ex-18 TFA salt is a complex macrocyclic molecule. It features a central benzimidazole core substituted with a fluorine atom. The macrocycle is formed by a chain of amide and ether linkages. Key substituents include a 4-methoxyphenyl group, a 1-hydroxyethyl group, and a 4-phenylbutyl group. The molecule is terminated with a trimethylammonium trifluoroacetate salt pair.	1539.75
Ex-19 TFA salt	 The chemical structure of Ex-19 TFA salt is a complex macrocyclic molecule, similar to Ex-18 but with a different ring closure. It features a central benzimidazole core substituted with a fluorine atom. The macrocycle is formed by a chain of amide and ether linkages. Key substituents include a 4-methoxyphenyl group, a 1-hydroxyethyl group, and a 4-phenylbutyl group. The molecule is terminated with a trimethylammonium trifluoroacetate salt pair.	1692.91
Ex-20 TFA salt	 The chemical structure of Ex-20 TFA salt is a complex macrocyclic molecule, similar to Ex-18 and Ex-19 but with a different ring closure. It features a central benzimidazole core substituted with a fluorine atom. The macrocycle is formed by a chain of amide and ether linkages. Key substituents include a 4-methoxyphenyl group, a 1-hydroxyethyl group, and a 4-phenylbutyl group. The molecule is terminated with a trimethylammonium trifluoroacetate salt pair.	1553.62

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-21 Cl salt	 The chemical structure of Ex-21 Cl salt is a complex molecule featuring a central benzimidazole core. It is substituted with a 4-fluorophenyl group, a 4-methoxyphenyl group, and a 4-(4-methoxyphenyl)pyrrolidin-1-yl group. The molecule also contains a 4-(4-methoxyphenyl)pyrrolidin-1-yl group and a 4-(4-methoxyphenyl)pyrrolidin-1-yl group. The structure is shown as a chloride salt, with a Cl⁻ ion indicated.	1566.91
Ex-22 TFA salt	 The chemical structure of Ex-22 TFA salt is a complex molecule featuring a central benzimidazole core. It is substituted with a 4-fluorophenyl group, a 4-methoxyphenyl group, and a 4-(4-methoxyphenyl)pyrrolidin-1-yl group. The molecule also contains a 4-(4-methoxyphenyl)pyrrolidin-1-yl group and a 4-(4-methoxyphenyl)pyrrolidin-1-yl group. The structure is shown as a trifluoroacetate salt, with a CF₃COO⁻ ion indicated.	1596.66

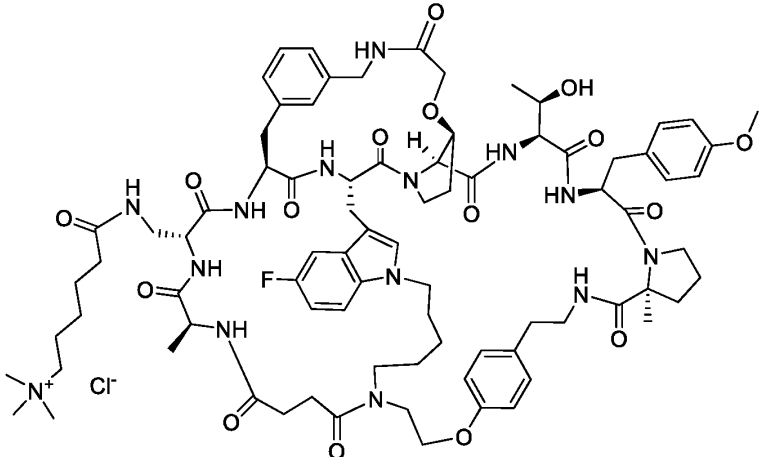
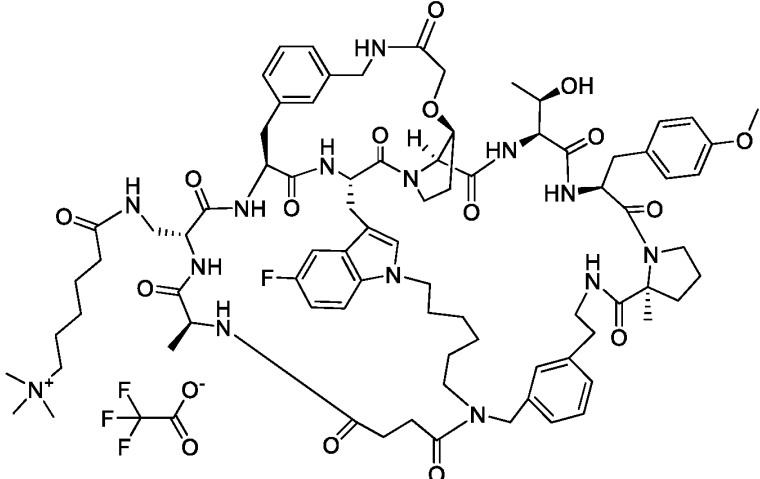
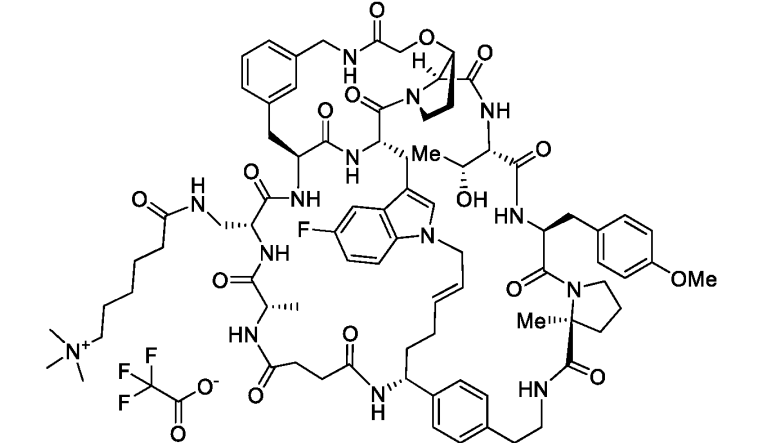
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-23		1656.57
Ex-24 TFA salt		1810.06 (half mass 848.64)

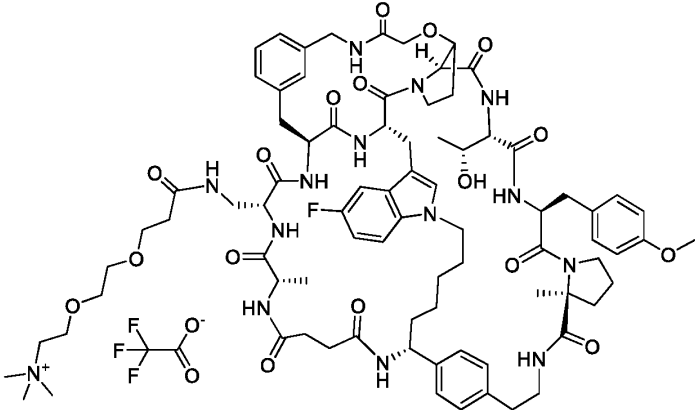
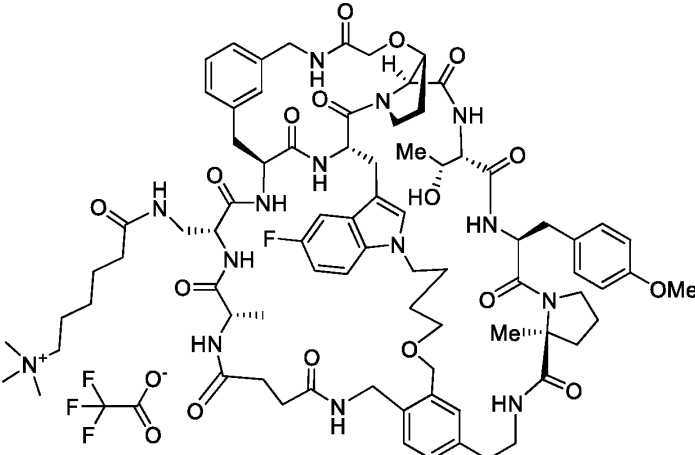
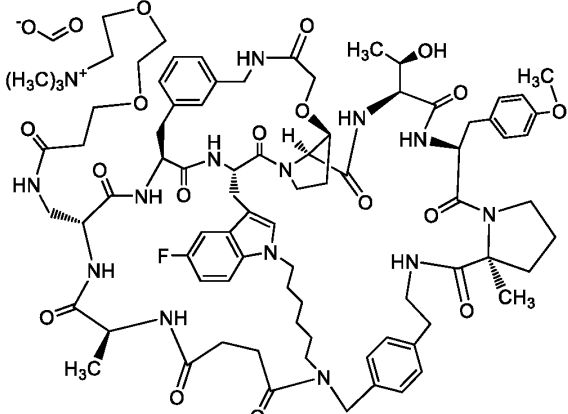
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-25 TFA salt		1550.6
Ex-25 Cl Salt		1550.6
Ex-26		1599.19

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-27		1628.74
Ex-28 TFA salt		1568.63
Ex-29 TFA salt		1566.60

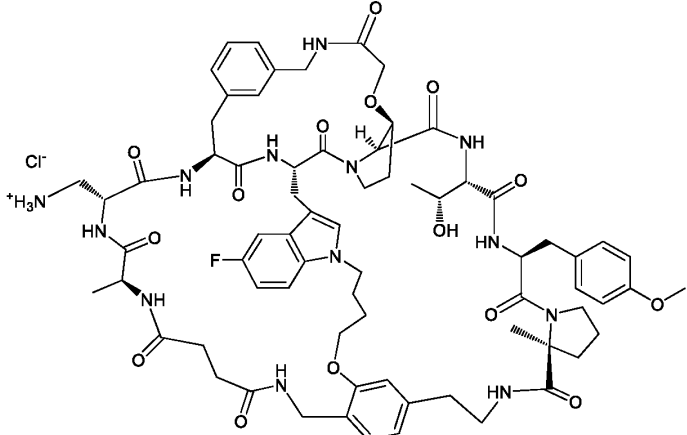
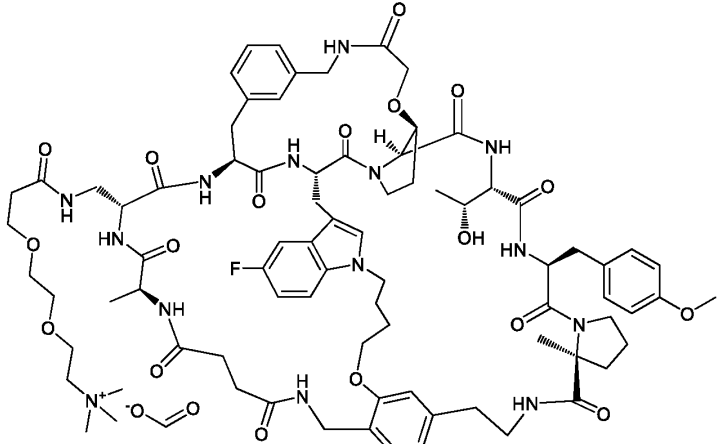
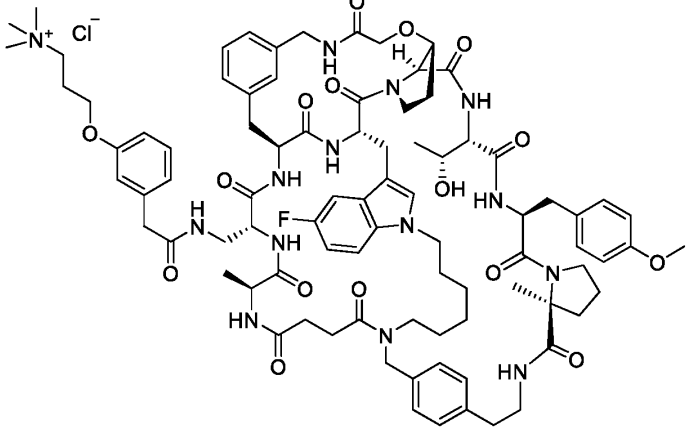
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-31 Cl salt		1552.85
Ex-35 Cl salt		1533.40
Ex-36 Cl salt		1520.60

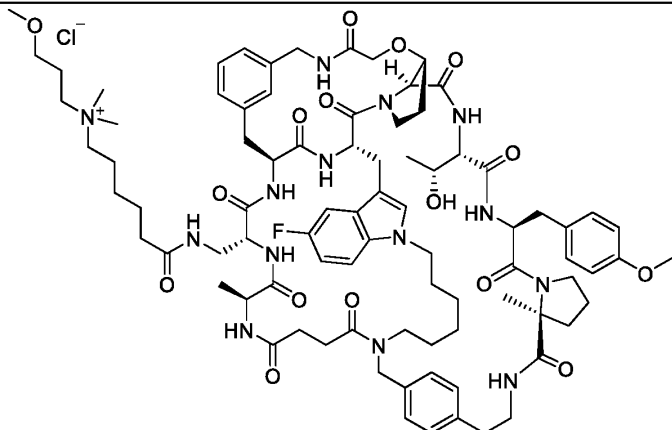
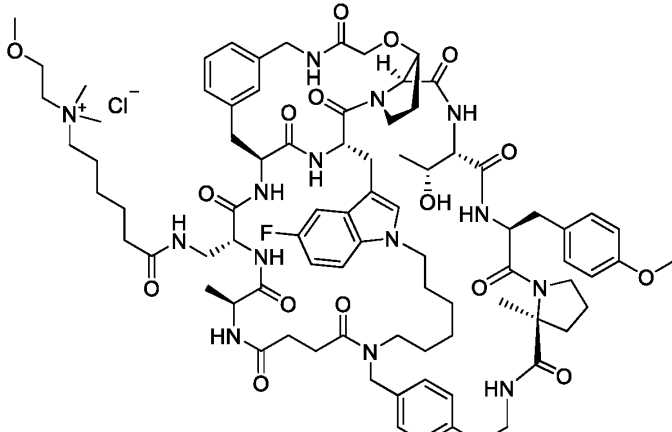
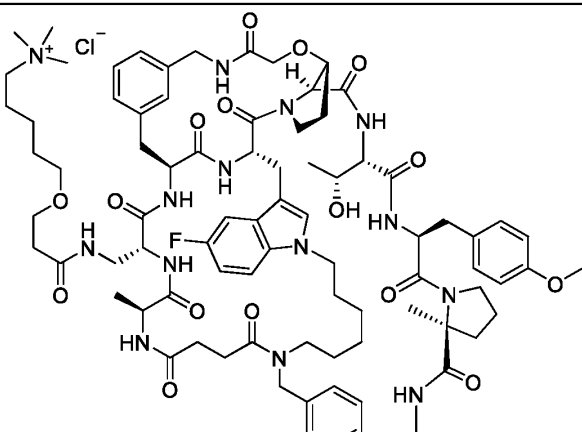
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-38 TFA salt		1536.21
Ex-39 TFA salt		1551.21
Ex-40 Cl salt		1538.60

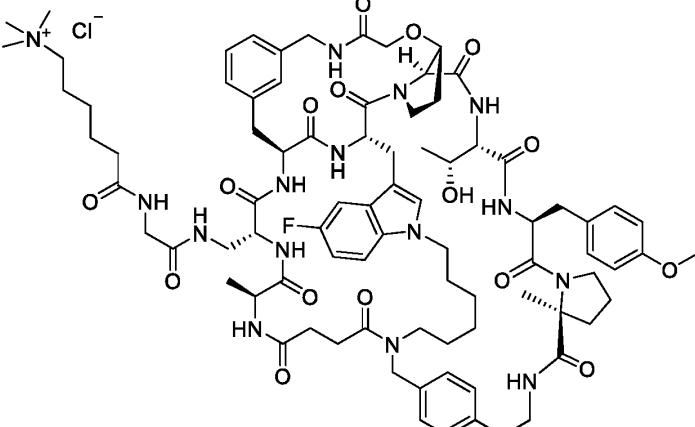
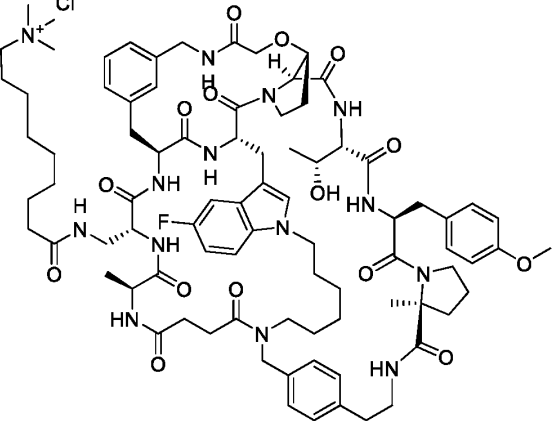
Ex- No/	Structure	LC/MS: (M) ⁺
Ex-41 Cl salt		1566.40
Ex-44 TFA salt		1549.49
Ex-47 TFA salt		1534.27

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-48 TFA salt		1582.20
Ex-49 TFA salt		1551.21
Ex-50 ACOH salt		1596.3

[illegible]

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-54 Cl salt	 The chemical structure of Ex-54 Cl salt is a complex macrocyclic molecule. It features a central indole ring system substituted with a fluorine atom and a long chain containing an ether linkage. The macrocycle is formed by several amide bonds and includes a cyclohexane ring with a hydroxyl group. A side chain contains a 4-methoxyphenyl group. The molecule is shown as a chloride salt with a protonated amine group (^+H_3N) and a chloride ion (Cl^-).	1383.44
Ex-55 ACOH salt	 The chemical structure of Ex-55 ACOH salt is a complex macrocyclic molecule, similar to Ex-54. It features a central indole ring system substituted with a fluorine atom and a long chain containing an ether linkage. The macrocycle is formed by several amide bonds and includes a cyclohexane ring with a hydroxyl group. A side chain contains a 4-methoxyphenyl group. The molecule is shown as an acetate salt with a quaternary ammonium cation (N^+) and an acetate anion (CH_3COO^-).	1583.69
Ex-56 Cl salt	 The chemical structure of Ex-56 Cl salt is a complex macrocyclic molecule, similar to Ex-54. It features a central indole ring system substituted with a fluorine atom and a long chain containing an ether linkage. The macrocycle is formed by several amide bonds and includes a cyclohexane ring with a hydroxyl group. A side chain contains a 4-methoxyphenyl group. The molecule is shown as a chloride salt with a quaternary ammonium cation (N^+) and a chloride ion (Cl^-).	1628.95

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-57 Cl salt		1608.95
Ex-58 Cl salt		1594.93
Ex-59 Cl salt		1593.7

Ex- No/	Structure	LC/MS: (M) ⁺
Ex-60 Cl salt		1607.4
Ex-61 Cl salt		1592.4

ACTIVITY DETERMINATION

Selected compounds of the invention were subjected to one or more of the following procedures to assay their activity for antagonism of PCSK9 activity.

The following is a description of the assays used to determine activity of compounds of the invention, and any comparator compounds reported, toward PCSK9 antagonism.

Biotinylated PCSK9 was obtained commercially.

LDLR TR-FRET

The PCSK9 TR-FRET assay measures the interaction between PCSK9 and LDLR. A solution containing 40 nM biotinylated PCSK9 + 10 nM Lance ULight Streptavidin is made in

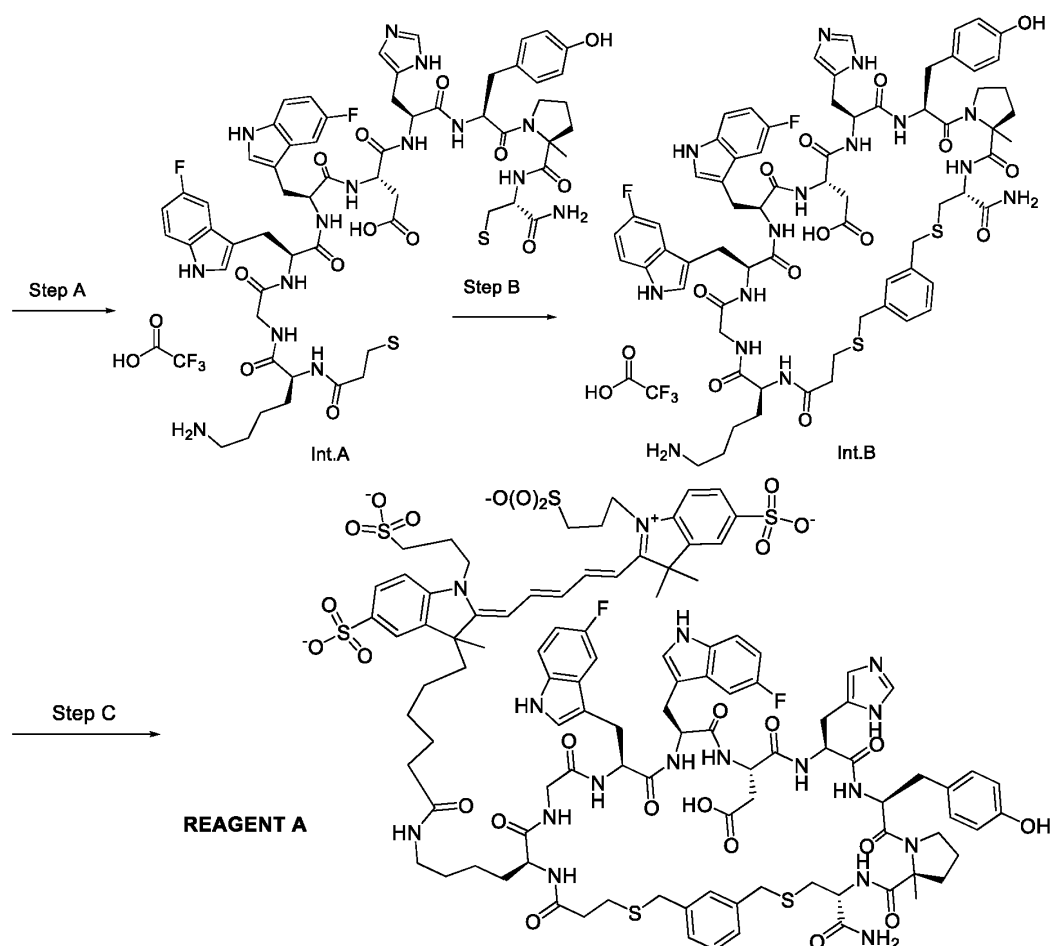
50 mM HEPES pH 7.4, 0.15 M NaCl, 5 mM CaCl₂, 0.01% BSA, and 0.01% Surfactant P20. A separate solution containing 40 nM rhLDLR-6xHis + 10 nM Eu-W1024 anti-6xHis is made in the same buffer system. An Echo is used to transfer 0.750ul of compound to an assay plate followed by the addition of 15ul of PCSK9+Ulight and 15ul of LDLR+Eu. The final assay
5 volume is 30.750ul containing 20nM PCSK9, 5nM Ulight, 20nM LDLR, and 5nM Eu. The reaction is incubated at room temperature for at least two hours prior to fluorescence measurements using an Envision Multilabel Reader. IC₅₀ values are determined by fitting data to a sigmoidal dose-response curve using nonlinear regression. Counts (B-counts) of the europium-labeled LDLR are followed to observe if compounds are adversely affecting
10 LDLR. A fall off of the B-counts is likely indicates a false positive of inhibition.

Alexa FRET Standard TR-FRET

The PCSK9 Alexa FRET Standard assay measures the interaction between PCSK9 and an AlexaFluor647 (AF) tagged cyclic peptide, Reagent A ($K_D = 83\text{nM}$). A solution containing 1 nM biotinylated PCSK9 + 2.5 nM Lance Streptavidin Europium (Strep-Eu) is
15 made in 50 mM HEPES pH 7.4, 0.15 M NaCl, 5 mM CaCl₂, 0.01% BSA, and 0.01% Surfactant P20. A separate solution containing 40 nM of the AlexaFluor tagged cyclic peptide is made in the same buffer system. An Echo is used to transfer 0.750ul of compound to an assay plate followed by the addition of 15ul of PCSK9+Strep-Eu and 15ul of AF peptide. The final assay volume is 30.750ul containing 0.5nM PCSK9, 1.25nM Strep-Eu, and 20nM AF
20 cyclic peptide. The reaction is incubated at room temperature for at least two hours prior to fluorescence measurements using an Envision Multilabel Reader. IC₅₀ values are determined by fitting data to a sigmoidal dose-response curve using nonlinear regression. K_i is then calculated from the IC₅₀ and the K_D of AF cyclic peptide. Counts (B-counts) of the europium-labeled PCSK9 are followed to observe if compounds are adversely PCSK9. A fall

off of the B-counts likely indicates a false positive of inhibition. Data from this procedure is reported as "A = 'numerical value' (nanomolar)"

Reagent A was prepared in accordance with the following method:



5

Step A – Synthesis of Intermediate Compound *Int-A*

The peptide was synthesized on a 0.250 mmol scale on CEM Liberty Blue, Microwave synthesizer using Fmoc/*t*Bu chemistry on PS Rink-Amide MBHA resin, 0.32 mmol g⁻¹. The assembly was performed using single-couplings using 4eq of Fmoc protected amino acid 0.2M in DMF, 4eq of 0.5M HATU in DMF, 4eq of 2M DIPEA (double coupling for Tyr). Fmoc deprotection cycles were performed using 20% (V/V) piperidine in DMF.

The sequence of Fmoc protected amino acids and building blocks used are:

1. N-(((9H-fluoren-9-yl)methoxy)carbonyl)-S-trityl-L-cysteine
2. (S)-1-(((9H-fluoren-9-yl)methoxy)carbonyl)amino)-2-methylpyrrolidine-2-carboxylic acid
3. (((9H-fluoren-9-yl)methoxy)carbonyl)-L-tyrosine
- 5 4. N-(((9H-fluoren-9-yl)methoxy)carbonyl)-N-trityl-L-histidine
5. (S)-2-(((9H-fluoren-9-yl)methoxy)carbonyl)amino)-4-(tert-butoxy)-4-oxobutanoic acid
6. (S)-2-(((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(5-fluoro-1H-indol-3-yl)propanoic acid
7. (S)-2-(((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(5-fluoro-1H-indol-3-yl)propanoic acid
- 10 8. (((9H-fluoren-9-yl)methoxy)carbonyl)glycine
9. N²-(((9H-fluoren-9-yl)methoxy)carbonyl)-N⁶-(tert-butoxycarbonyl)-L-lysine
10. 3-(tritylthio)propanoic acid

At the end of the assembly, the resin was washed with DMF, MeOH, DCM, Et₂O. The peptide was cleaved from solid support using 50 ml of TFA solution (v/v) (91% TFA, 5% H₂O, 4% TIPS) for approximately 1.5 hours, at room temperature. The resin was filtered, washed with TFA and solution concentrated to dryness and lyophilized. Lyophilization afforded *Intermediate Compound Int. A* (399mg), which was used as crude in the next step. LCMS anal. calcd. C₆₁H₇₅F₂N₁₅O₁₃S₂: 1328.48, found: 1328.2 (M+1)⁺

20 *Step B – Synthesis of Intermediate Compound Int-B:* as described for reagent B

Purified by RP-HPLC (Waters Deltapak C4, double cartridge, 40x100 mm, 15μm, 300A; 15% to 35% ACN/water + 0.1% TFA modifier over 20 min). Collected fractions lyophilized to afford 35mg of *Intermediate Compound Int-B*. LCMS anal. calcd. for C₆₉H₈₁F₂N₁₅O₁₃S₂: 1430.62; found: 1430.9 (M+1)⁺

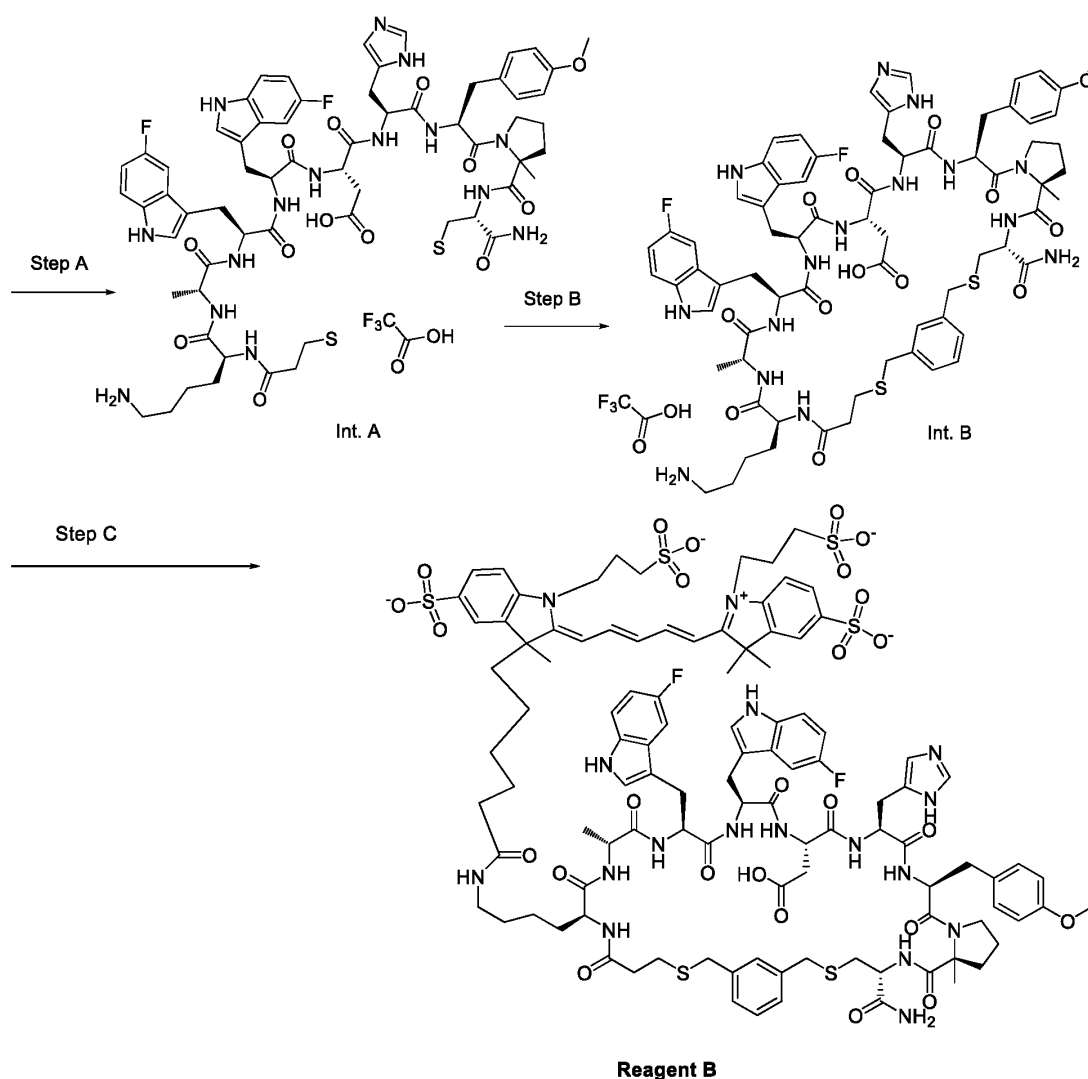
25 *Step C – Synthesis of Compound ReagentA:* as described for reagent B

LCMS anal. calcd. for C₁₀₅H₁₂₂F₂N₁₇O₂₆S₆³⁻: 2268.58; 1135.8 (M+2)²⁺

Alexa FRET Plus TR-FRET

The PCSK9 Alexa FRET Plus assay measures the interaction between PCSK9 and an AlexaFluor647 (AF) tagged cyclic peptide, Reagent B (K_D = 35nM). A solution
5 containing 1 nM biotinylated PCSK9 + 2.5 nM Lance Streptavidin Europium (Strep-Eu) is made in 50 mM HEPES pH 7.4, 0.15 M NaCl, 5 mM CaCl₂, 0.01% BSA, and 0.01% Surfactant P20. A separate solution containing 1920 nM of the AlexaFluor tagged cyclic peptide is made in the same buffer system. An Echo is used to transfer 0.075ul of compound plus 0.675ul of DMSO to each well of an assay plate followed by the addition of
10 15ul of PCSK9+Strep-Eu and 15ul of AF peptide. The final assay volume is 30.750ul containing 0.5nM PCSK9, 1.25nM Strep-Eu, and 960nM AF cyclic peptide. The reaction is incubated at room temperature for at least two hours prior to fluorescence measurements using an Envision Multilabel Reader. IC₅₀ values are determined by fitting data to a sigmoidal dose-response curve using nonlinear regression. K_i is then calculated from the
15 IC₅₀ and the K_D of AF cyclic peptide. Counts (B-counts) of the europium-labeled PCSK9 are followed to observe if compounds are adversely affecting PCSK9. A fall off of the B-counts is likely indicates a false positive of inhibition. Data from this procedure is reported as "P= 'numerical value' (nanomolar)"

Reagent B was prepared by the following procedure.



Step A – Synthesis of Intermediate Compound *Int-A*

The peptide was synthesized on a 0.250 mmol scale on CEM Liberty Blue, Microwave synthesizer using Fmoc/*t*Bu chemistry on PS Rink-Amide MBHA resin, 0.32 mmol g⁻¹. The assembly was performed using single-couplings using 4eq of Fmoc protected amino acid 0.2M in DMF, 4eq of 1M Oxyme in DMF, 4eq of 0.5M *N,N*-diisopropylcarbodiimide (DIC) (double coupling for Y01). Fmoc deprotection cycles were performed using 20% (V/V) piperidine in DMF.

The sequence of Fmoc protected amino acids and building blocks used are:

1. N-(((9H-fluoren-9-yl)methoxy)carbonyl)-S-trityl-L-cysteine

2. (S)-1((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-2-methylpyrrolidine-2-carboxylic acid
3. (S)-2((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(4-methoxyphenyl)propanoic acid
- 5 4. N-((((9H-fluoren-9-yl)methoxy)carbonyl)-N-trityl-L-histidine
5. (S)-2((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-4-(tert-butoxy)-4-oxobutanoic acid
6. (S)-2((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(5-fluoro-1H-indol-3-yl)propanoic acid
7. (S)-2((((9H-fluoren-9-yl)methoxy)carbonyl)amino)-3-(5-fluoro-1H-indol-3-yl)propanoic acid
- 10 8. (((9H-fluoren-9-yl)methoxy)carbonyl)-D-alanine
9. N²-((((9H-fluoren-9-yl)methoxy)carbonyl)-N⁶-(tert-butoxycarbonyl)-L-lysine
10. 3-(tritylthio)propanoic acid

At the end of the assembly, the resin was washed with DMF, MeOH, DCM, Et₂O. The peptide was cleaved from solid support using 50 ml of TFA solution (v/v) (91% TFA, 5% H₂O, 4% TIPS) for approximately 1.5 hours, at room temperature. The resin was filtered, washed with TFA and solution concentrated to dryness and lyophilized. Lyophilization afforded *Intermediate Compound Int. A* (300mg), which was used as crude in the next step. LCMS anal. calcd. C₆₃H₇₉F₂N₁₅O₁₃S₂: 1356.53, found: 1356.9 (M+1)⁺

20 Step B – Synthesis of Intermediate Compound *Int-B*

Crude *Int-A* (0.22 mmol) was redissolved in 24ml of DMF. 6ml of 1M aqueous solution of sodium bicarbonate was added to raise the pH to 7. Then 0.26 mmol of 1,3-bis(bromomethyl)benzene (0.1M in DMF) were added dropwise. Reaction was left under stirring at room temperature for 20 min, quenched with TFA (pH to 3-4) and then concentrated in vacuo to provide crude *Int-B*, which was purified by RP-HPLC (Waters

XBridge, C18, 50x150 mm, 5 μ m, 130A; 25% to 40% ACN/water + 0.1% TFA modifier over 20 min). Collected fractions were lyophilized to afford 35mg of *Intermediate Compound Int-B*. LCMS anal. calcd. for C71H85F2N15O13S2: 1458.67; found: 1458.8 (M+1)⁺

Step C – Synthesis of Compound Reagent B

- 5 *Intermediate Compound Int-B* (15mg) was dissolved in 0.2ml of dry DMSO. Then 15mg of ALEXAFLUOR 647NHS Ester (A37566, Life technology) dissolved in 1.5ml of dry DMSO were added. 20 μ L of dry DIPEA were added. Reaction was left under stirring at room temperature for 12h under Nitrogen atmosphere in the dark. Quenched with TFA (pH to 3-4) and purified by RP-HPLC (Dr Maish, Reprosil Gold C18, 250x20 mm, 120 Å, 10 μ m; 20%
10 to 35% of 0.1% TFA in ACN/0.1% TFA in H₂O, over 20min, then 35% to 40% over 5min at 20 mL/min flow rate). Collected fractions were lyophilized to afford 16.1 mg of *Compound Reagent B*. LCMS anal. for C107H126F2N17O26S6³⁻: 2296.64; found: 1150.6 (M+2)²⁺

Activity data obtained by one or both of the above-described procedures is reported for selected example compounds of the invention in the following format:

- 15 Example No.: A (standard TR Fret) = 'numerical value'; P (Alexa Fret plus standard TR Fret) = 'numerical value' /, note that all values reported are nanomolar.

Alexa FRET Ultra TR-FRET

- The PCSK9 Alexa FRET Ultra assay measures the interaction between PCSK9 and an AlexaFluor647 (AF) tagged cyclic peptide, Reagent B (K_D = 0.99nM). A solution
20 containing 1 nM biotinylated PCSK9 + 2.5 nM Lance Streptavidin Europium (Strep-Eu) is made in 50 mM HEPES pH 7.4, 0.15 M NaCl, 5 mM CaCl₂, 0.01% BSA, and 0.01% Surfactant P20. A separate solution containing 1920 nM of the AlexaFluor tagged cyclic peptide is made in the same buffer system. An Echo is used to transfer 0.015 μ L of compound plus 0.735 μ L of DMSO to each well of an assay plate followed by the addition of
25 15 μ L of PCSK9+Strep-Eu and 15 μ L of AF peptide. The final assay volume is 30.750 μ L

containing 0.5 nM PCSK9, 1.25 nM Strep-Eu, and 960 nM AF cyclic peptide. The reaction is incubated at room temperature for at least two hours prior to fluorescence measurements using an Envision Multilabel Reader. IC50 values are determined by fitting data to a sigmoidal dose-response curve using nonlinear regression. Ki is then calculated from the IC50 and the K_D of AF cyclic peptide. Counts (B-counts) of the europium-labeled PCSK9 are followed to observe if compounds are adversely affecting PCSK9. A fall off of the B-counts is likely indicates a false positive of inhibition. Data from this procedure is reported as "Ki Ultra = 'numerical value' (data reported is nanomolar)"

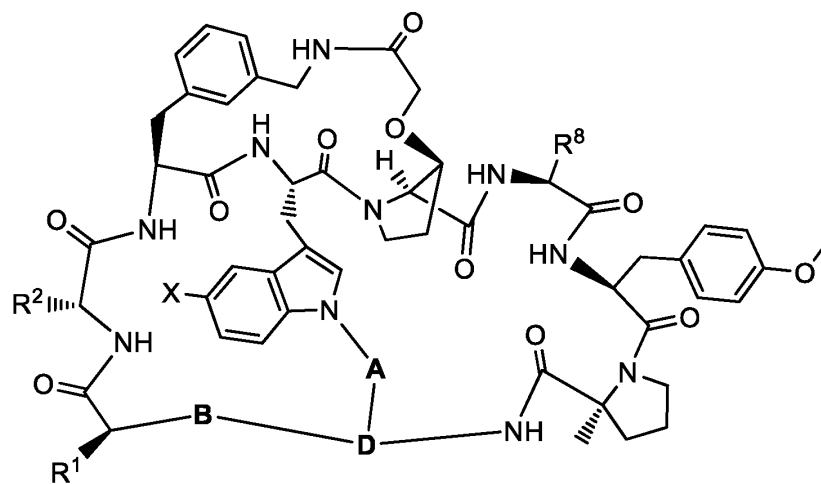
The following compounds were assessed, as shown in Table 2, using the protocol described above with the results shown:

Ex-01 Ki Plus = < 0.00558, Ki Ultra = 0.0046 / **Ex-02** Ki Plus = 0.00558, Ki Ultra = 0.005933/ **Ex-03** Ki Plus = 0.02535, Ki Ultra = 0.06803/ **Ex-04** Ki Plus ≤ 0.00558, Ki Ultra = 0.004711/ **Ex-05** Ki Plus = 0.009621, Ki Ultra = 0.03296/ **Ex-06** Ki Plus = 0.00568, Ki Ultra = 0.003424/ **Ex-07** Ki Plus = 0.05914, Ki Ultra = 0.06753/ **Ex-08** Ki Plus = 0.01574, Ki Ultra = 0.06832/ **Ex-09** Ki Plus = 0.09189, Ki Ultra = 0.247/ **Ex-10** Ki Plus = 0.005743, Ki Ultra = 0.02489/ **Ex-11** Ki Plus = 0.04334, Ki Ultra = 0.2067/ **Ex-12** Ki Plus = 0.01448, Ki Ultra = 0.02247/ **Ex-13** Ki Plus = 0.1454, Ki Ultra = 0.4772/ **Ex-14** Ki Plus = 0.01605, Ki Ultra = 0.02099/ **Ex-15** Ki Plus = 0.1027, Ki Ultra = 0.2601/ **Ex-16** Ki Plus = 0.01423, Ki Ultra = 0.05141/ **Ex-17** Ki Plus ≤ 0.00558, Ki Ultra = 0.0028/ **Ex-18** Ki Plus = 0.03356, Ki Ultra = 0.1183/ **Ex-19** Ki Plus = 0.01662, Ki Ultra = 0.01204/ **Ex-20** Ki Plus = 0.01303, Ki Ultra = 0.01711/ **Ex-21** Ki Plus = 0.005692, Ki Ultra = 0.001264/ **Ex-22** Ki Plus = 0.00926, Ki Ultra = 0.01519/ **Ex-23** Ki Plus = 0.00938, Ki Ultra = 0.00239/ **Ex-24** Ki Plus = 0.00812, Ki Ultra = 0.00767/ **Ex-25** Ki Plus = 0.01127, Ki Ultra = 0.00463/ **Ex-26** Ki Plus ≤ 0.00558, Ki Ultra = 0.002754/ **Ex-27** Ki Plus ≤ 0.00558, Ki Ultra = 0.00301/ **Ex-28** Ki Plus ≤ 0.00558, Ki Ultra = 0.00078/ **Ex-29** Ki Plus = 0.00981, Ki Ultra = 0.00614/ **Ex-31** Ki Plus < 0.00558, Ki Ultra = 0.00074/ **Ex-35** Ki Plus = 0.04652, Ki Ultra = 0.08434/ **Ex-36** Ki Plus = 0.00762, Ki Ultra = 0.00507/ **Ex-38** Ki Plus = 0.00904, Ki

Ultra = 0.01416/ **Ex-39** Ki Plus = 0.00716, Ki Ultra = 0.00414/ **Ex-40** Ki Plus
= 0.30800, Ki Ultra = 0.86010/ **Ex-41** Ki Plus = 0.00697, Ki Ultra = 0.00628/
Ex-44 Ki Plus = 0.01445, Ki Ultra = 0.02194/ **Ex-47** Ki Plus = 0.01474, Ki
Ultra = 0.01193/ **Ex-48** Ki Plus = 0.01169, Ki Ultra = 0.01545/ **Ex-49** Ki Plus
= 0.00716, Ki Ultra = 0.00414/ **Ex-50** Ki Standard <1.26, Ki Plus = 0.01052,
Ki Ultra = 0.00443/ **Ex-51** Ki Standard <1.26, Ki Plus < 0.00558, Ki Ultra =
0.00597/ **Ex-52** Ki Standard <1.26, Ki Plus < 0.00558, Ki Ultra = 0.00359/
Ex-53 Ki Standard <1.26, Ki Plus = 0.09629, Ki Ultra = 0.21500/ **Ex-54** Ki
Standard <1.26, Ki Plus = 0.36720, Ki Ultra = 0.48390/ **Ex-55** Ki Standard
<1.26, Ki Plus = 0.07240, Ki Ultra = 0.23800/ **Ex-56** Ki Standard <1.257, Ki
Plus = 0.02237, Ki Ultra = 0.00481/ **Ex-57** Ki Standard <1.257, Ki Plus <
0.00558, Ki Ultra = 0.00162/ **Ex-58** Ki Standard <1.257, Ki Plus = 0.00773,
Ki Ultra = 0.00196/ **Ex-59** Ki Plus = 0.00788, Ki Ultra = 0.004959/ **Ex-60** Ki
Plus = 0.006515, Ki Ultra = 0.005312 / **Ex-61** Ki Plus = 0.00747, Ki Ultra =
0.006543.

What is Claimed is:

- 1) A compound of Formula I:



Formula I,

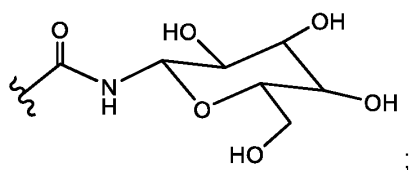
wherein:

X is H F, Cl or Br;

R¹ is selected from:

- (a) -H; or
- (b) $-(CH_2)_z-R^{14A}$, wherein: z is 1-6, and R^{14A} is:
 - (i) -H;
 - (ii) $-NH_2$;
 - (iii) $-N^+H_3$;
 - (iv) $-N^+(H_3C)_3$;
 - (v) $-NH-C(O)-[(CH_2)_2-O]_2-(CH_2)_2R^{14B}$ wherein R^{14B} is: $-NH_2$; $-N^+H_3$; $-N(CH_3)_2$; or $-N^+(CH_3)_3$;
 - (vi) $-NH-C(O)-[(CH_2)_{y12}-O]_2-(CH_2)_{y13}R^{14B}$ wherein: y₁₂ and y₁₃ are not both 2 and are independently 2 to 4; and R^{14B} is: $-NH_2$; $-N^+H_3$; $-N(CH_3)_2$; or $-N^+(CH_3)_3$;

- (vii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y = 1$ to 6 and $\text{R}^{14\text{C}}$ is $-\text{O}-(\text{CH}_2)_{3-4}-\text{N}^+(\text{CH}_3)_3$; and
- (viii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y = 1$ to 6 and $\text{R}^{14\text{C}}$ is:
- (ai) $-\text{O}-(\text{CH}_2)_2-\text{N}^+(\text{CH}_3)_3$;
 - (aii) $-\text{N}^+(\text{CH}_3)_3$; or
 - (aiii) a moiety of the formula:

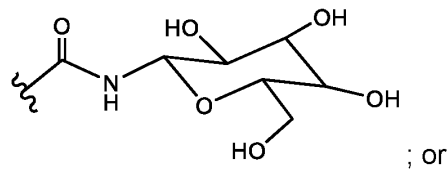


R^2 is selected from:

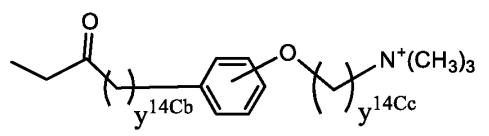
- (a) $-\text{H}$; and
- (b) $-(\text{CH}_2)_z-\text{R}^{14\text{A}}$, wherein: z is $1-6$, and $\text{R}^{14\text{A}}$ is selected from:
 - (i) $-\text{H}$;
 - (ii) $-\text{NH}_2$;
 - (iii) $-\text{N}^+\text{H}_3$;
 - (iv) $-\text{N}^+(\text{H}_3\text{C})_3$;
 - (v) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_2-\text{O}]_2-(\text{CH}_2)_2\text{R}^{14\text{B}}$ wherein $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
 - (vi) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_{y12}-\text{O}]_2-(\text{CH}_2)_{y13}\text{R}^{14\text{B}}$ wherein: $y12$ and $y13$ are not both 2 and are independently 2 to 4 ; and $\text{R}^{14\text{B}}$ is: $-\text{NH}_2$; $-\text{N}^+\text{H}_3$; $-\text{N}(\text{CH}_3)_2$; or $-\text{N}^+(\text{CH}_3)_3$;
 - (vii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y = 1$ to 6 and $\text{R}^{14\text{C}}$ is $-\text{O}-(\text{CH}_2)_{3-4}-\text{N}^+(\text{CH}_3)_3$; and
 - (viii) $-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_y\text{R}^{14\text{C}}$, wherein, $y = 1$ to 6 and $\text{R}^{14\text{C}}$ is:
 - (ai) $-\text{O}-(\text{CH}_2)_2-\text{N}^+(\text{CH}_3)_3$;

(aii) $-N^+(CH_3)_2R^{14ca}$, wherein R^{14ca} is $-CH_3$ or $-(CH_2)_{1-4}-OCH_3$;

(aiii) a moiety of the formula:

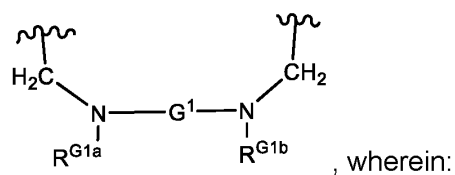


(aiv) a moiety of the formula:



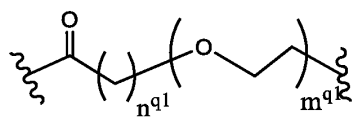
where R^{14Cb} and R^{14Cc} are 1 to 4; or

R^1 and R^2 may be bonded together to form a moiety of the formula:



G^1 , R^{G1a} and R^{G1b} are defined as follows:

(a) G^1 is a linker moiety of the formula:

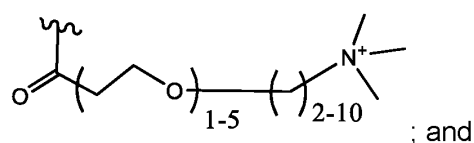


wherein n^{q1} is 1 to 6, m^{q1} is 0, 1 or 2 and together the value of n^{q1} and m^{q1} are selected such that the length of the linker moiety they define does not exceed a total of 8 carbon and/or oxygen atoms comprising the chain including the carbon atom in the chain that forms the carbonyl moiety;

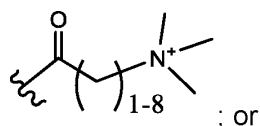
R^{G1a} is selected from: (i) $-H$; and (ii) alkyl of up to 4 carbon atoms; and

R^{G1b} is selected from:

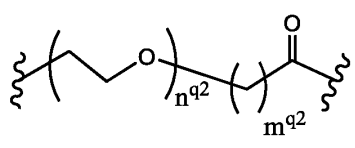
(i) a moiety of the formula:



(ii) a moiety of the formula:



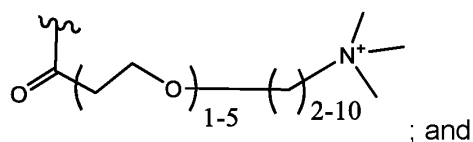
(b) G¹ is a linker moiety of the formula:



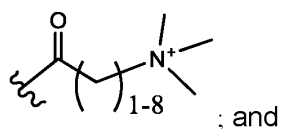
wherein n^{q2} is 0, 1 or 2, m^{q2} is 1 to 6, and together the value of n^{q2} and m^{q2} are selected such that the length of the linker moiety they define does not exceed a total of 8 carbon and/or oxygen atoms comprising the chain including the carbon atom in the chain that forms the carbonyl moiety;

R^{G1a} is selected from:

(i) a moiety of the formula:

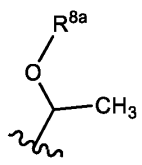


(ii) a moiety of the formula:



R^{G1b} is selected from: (i) -H; and (ii) alkyl of up to 4 carbon atoms;

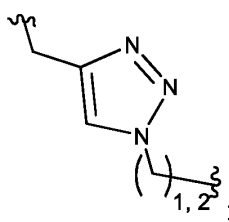
R⁸ is -CH₃ or a moiety of the formula:



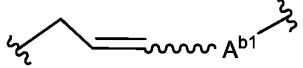
wherein R^{8a} is $-H$, or a linear, branched or cyclic alkyl of up to four carbon atoms;

A is selected from:

(a) a moiety of the formula:

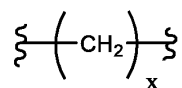


(b) $-CH_2-(CH_2)_y-CH_2-$, wherein y is 1 to 6;

(c) a moiety of the formula: 

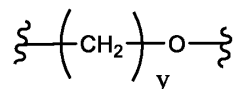
wherein A^{b1} is:

(i) a moiety of the formula:



wherein x is 1 to 6; or

(ii) a moiety of the formula:

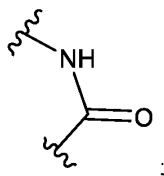


wherein y is 1 to 5;

(d) a moiety of the formula: $-CH_2-(CH_2)_m-O-(CH_2)_n-$, wherein $m = 1$ to 5, and $n = 0$ or 1 to 4;

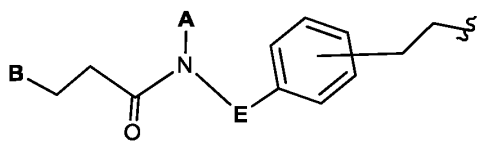
B is:

- (a) a bond;
- (b) $-(CH_2)_{1-2}-$; or
- (c) a moiety of the formula:



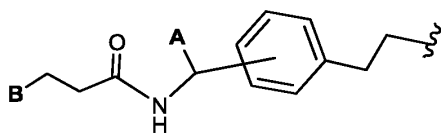
D is:

- (a) a moiety of the Formula:



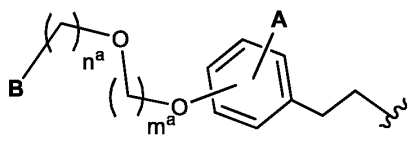
wherein **E** is $-CH_2-$ or $-(CH_2)_{2-4}-O-$, and **A** and **B** are as defined above;

- (b) a moiety of the formula:



wherein **A** and **B** are as defined above;

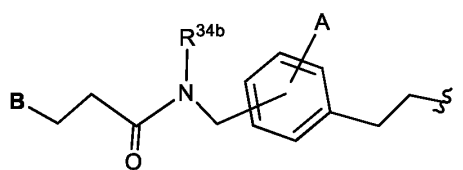
- (c) a moiety of the formula:



wherein n^a is 1, 2, or 3, m^a is 2, 3, or 4, and $n^a + m^a$ is ≥ 3 , and wherein **A** and

B are as defined above;

- (d) a moiety of the formula:

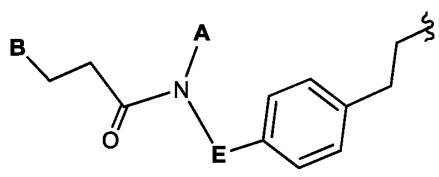


wherein, R^{34b} is $-H$ or a linear, branched or cyclic alkyl of up to four carbon atoms, and **A** and **B** are as defined above,

or a pharmaceutically acceptable salt of any thereof.

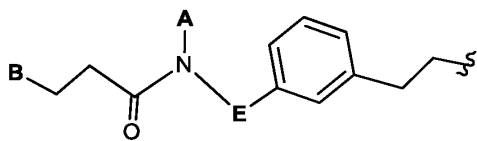
2. A compound according to Claim 1 of **Formula I**, wherein **X** is **F**, or a pharmaceutically acceptable salt of any thereof.

3. A compound of Claim 2 wherein **D** is a moiety of the formula:



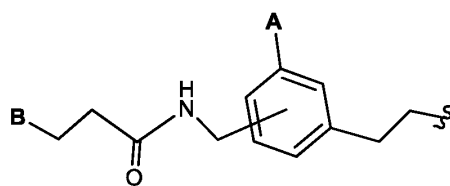
wherein, **E** is $-CH_2-$ or $-(CH_2)_2-O-$, or a pharmaceutically acceptable salt thereof.

4. A compound of Claim 2 wherein **D** is a moiety of the formula:



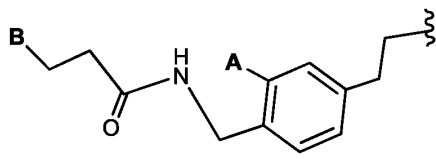
wherein, **E** is $-CH_2-$ or $-(CH_2)_2-O-$, or a pharmaceutically acceptable salt thereof.

5. A compound of Claim 2 wherein **D** is a moiety of the formula:



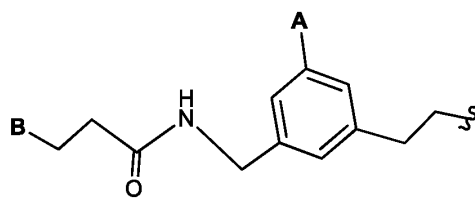
or a pharmaceutically acceptable salt thereof.

6. A compound of Claim 2 wherein **D** is a moiety of the formula:



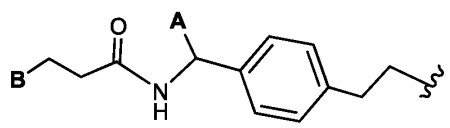
or a pharmaceutically acceptable salt thereof.

7. A compound of Claim 2 wherein **D** is a moiety of the formula:



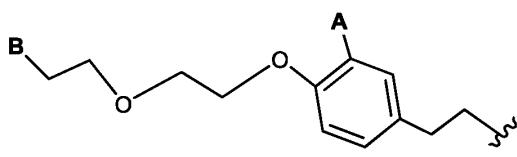
or a pharmaceutically acceptable salt thereof.

8. A compound of Claim 2 wherein **D** is a moiety of the formula:



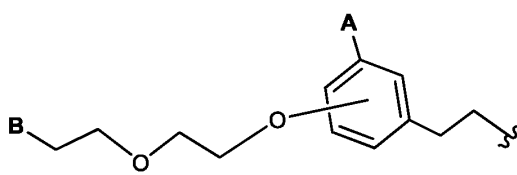
or a pharmaceutically acceptable salt thereof.

9. A compound of Claim 2 wherein **D** is a moiety of the formula:



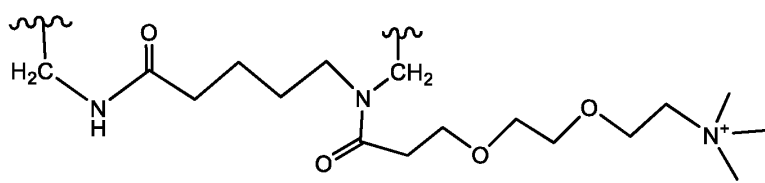
or a pharmaceutically acceptable salt thereof.

10. A compound of Claim 2 wherein **D** is a moiety of the formula:



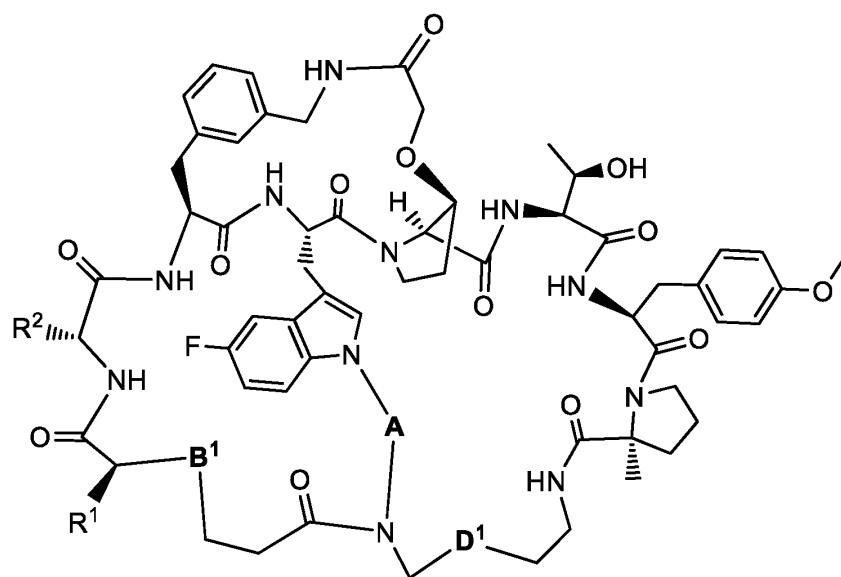
or a pharmaceutically acceptable salt thereof.

11. A compound of Claim 2 wherein **R**¹ and **R**² are joined together with a moiety of the formula:



such that together with the cyclopeptide to which **R**¹ and **R**² are attached they form acyclic structure,
or a pharmaceutically acceptable salt thereof.

12. A compound of Claim 2 having the structure of **Formula IIB**, or a pharmaceutically acceptable salt thereof:

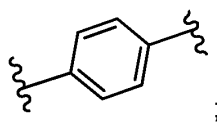


Formula IIB,

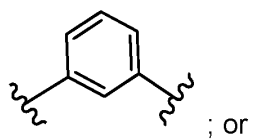
B¹ is $-(\text{CH}_2)_{0-2}-$; and

D¹ is:

- (a) a moiety of the formula:

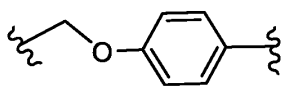


- (b) a moiety of the formula:

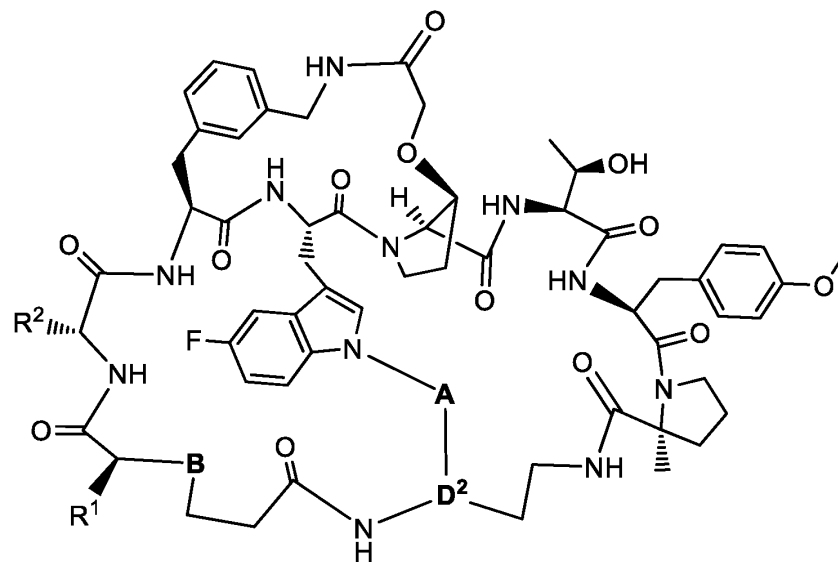


; or

- (c) a moiety of the formula:



13. A compound of Claim 2 having the structure of **Formula IIC**, or a pharmaceutically acceptable salt thereof:

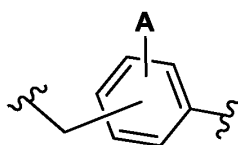


Formula IIC,

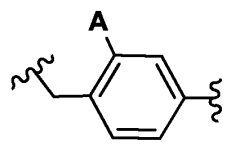
wherein

D² is:

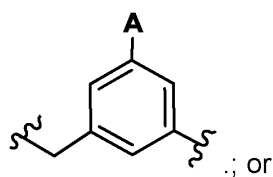
- (a) a moiety of the formula:



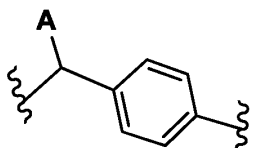
- (b) a moiety of the formula:



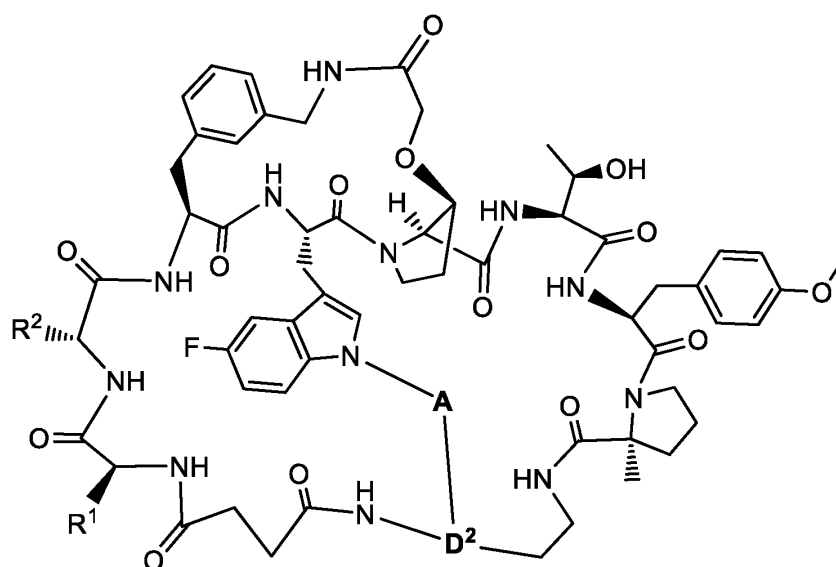
- (c) a moiety of the formula:



(d) a moiety of the formula:

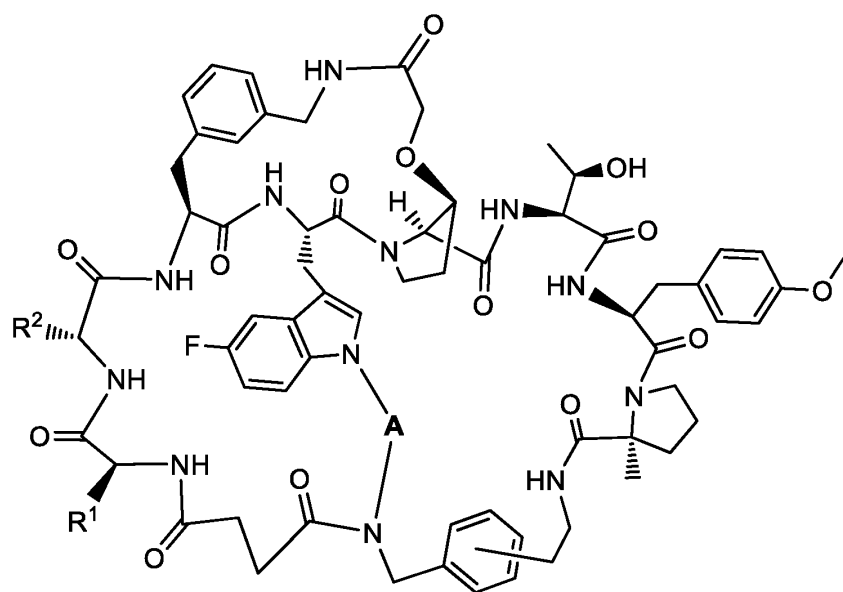


14. A Compound of Claim 2 having the structure of Formula IID, or a pharmaceutically acceptable salt thereof:



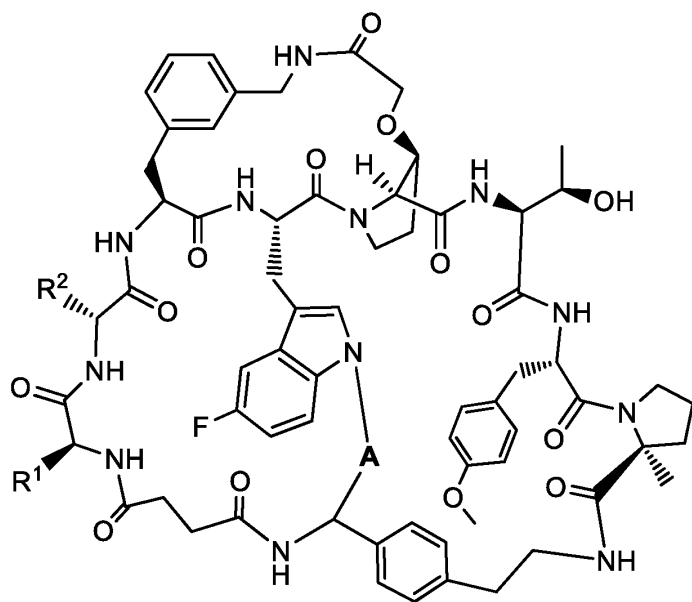
Formula IID.

15. A compound of Claim 2 having the structure of **Formula IIE**, or a pharmaceutically acceptable salt thereof:



Formula IIE.

16. A compound of Claim 2 having the structure of Formula IIF, or a pharmaceutically acceptable salt thereof:

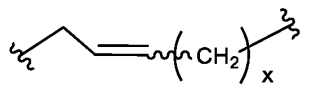


Formula IIF.

17. A compound of any of Claims 1 to 16, or a pharmaceutically acceptable salt thereof, wherein **A** is:

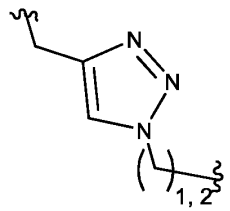
(a) $-(CH_2)_6$;

(b) a moiety of the formula:



wherein x is 1 to 3; or

(c) a moiety of the formula:



18. A compound of any of Claims 1 to 17, or a pharmaceutically acceptable salt thereof, wherein R^2 is:

(a) $-(CH_2)_z-R^{14A}$, wherein: z is 1-6, and R^{14A} is:

(a) $-H$;

(b) $-CH_3$;

(c) $-NH_2$;

(d) $-N^+H_3$;

(e) $-N^+(H_3C)_3$;

(f) $-NH-C(O)-[(CH_2)_{2-4}-O]_{2-4}-(CH_2)_{2-4}R^{14B}$ wherein R^{14B}

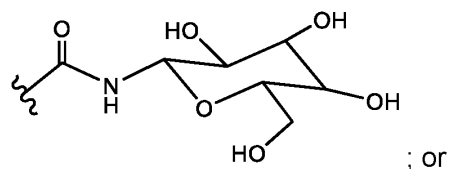
is: $-NH_2$; $-N^+H_3$; $-N(CH_3)_2$; or $-N^+(CH_3)_3$;

(g) $-NH-C(O)-[(CH_2)_yR^{14C}]$, wherein, y= 1 to 6 and R^{14C} is:

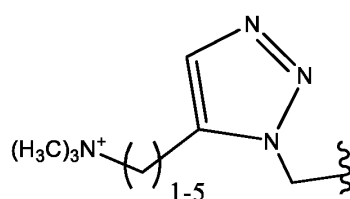
(ai) $-\text{O}-(\text{CH}_2)_{2-4}-\text{N}^+(\text{CH}_3)_3$;

(a ii) $-\text{N}^+(\text{CH}_3)_3$; or

(a iii) a moiety of the formula:



(b) a moiety of the formula



19. A compound of any of Claims 1 to 18, or a pharmaceutically acceptable salt thereof, wherein R^1 is selected from:

(a) $-\text{H}$;

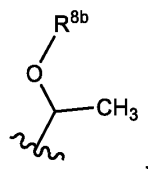
(b) $-(\text{CH}_2)_z-\text{R}^{14\text{A}}$, wherein: z is 1-6, and $\text{R}^{14\text{A}}$ is:

(i) $-\text{H}$;

(ii) $-\text{N}^+\text{H}_3$; or

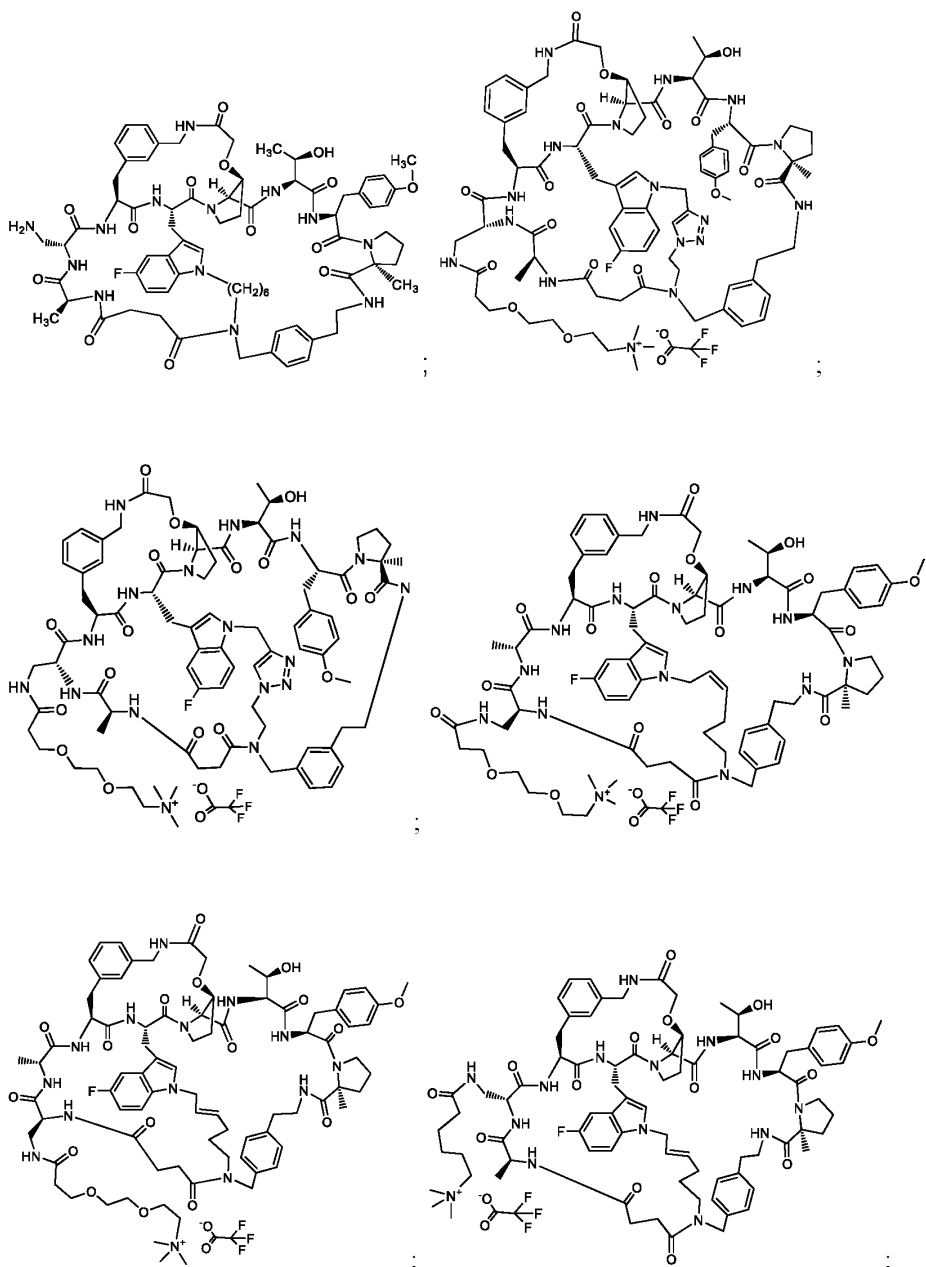
(iii) $-\text{NH}-\text{C}(\text{O})-[(\text{CH}_2)_2-\text{O}]_2-(\text{CH}_2)_2-\text{N}^+(\text{CH}_3)_3$.

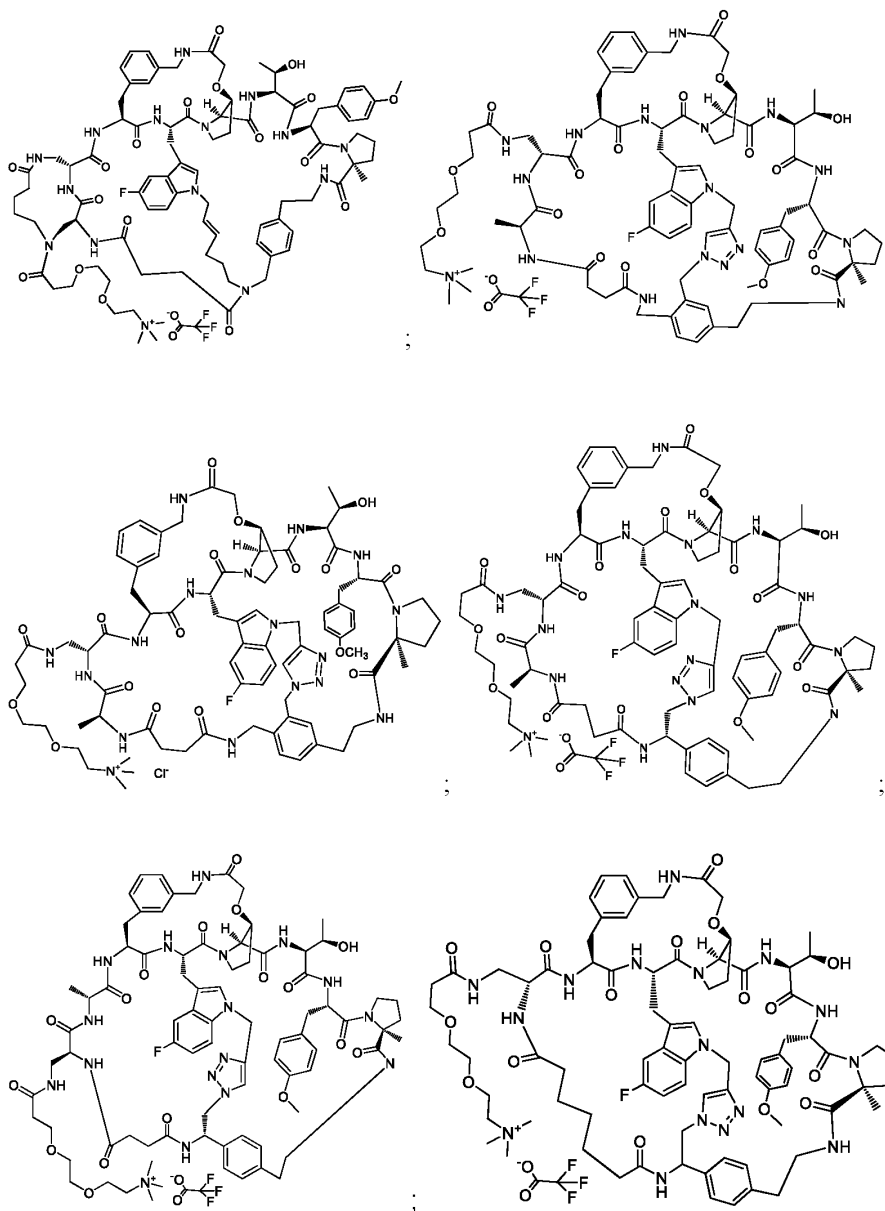
20. A compound of any of Claims 1 to 19, or a pharmaceutically acceptable salt thereof, wherein R^8 is a moiety of the formula:

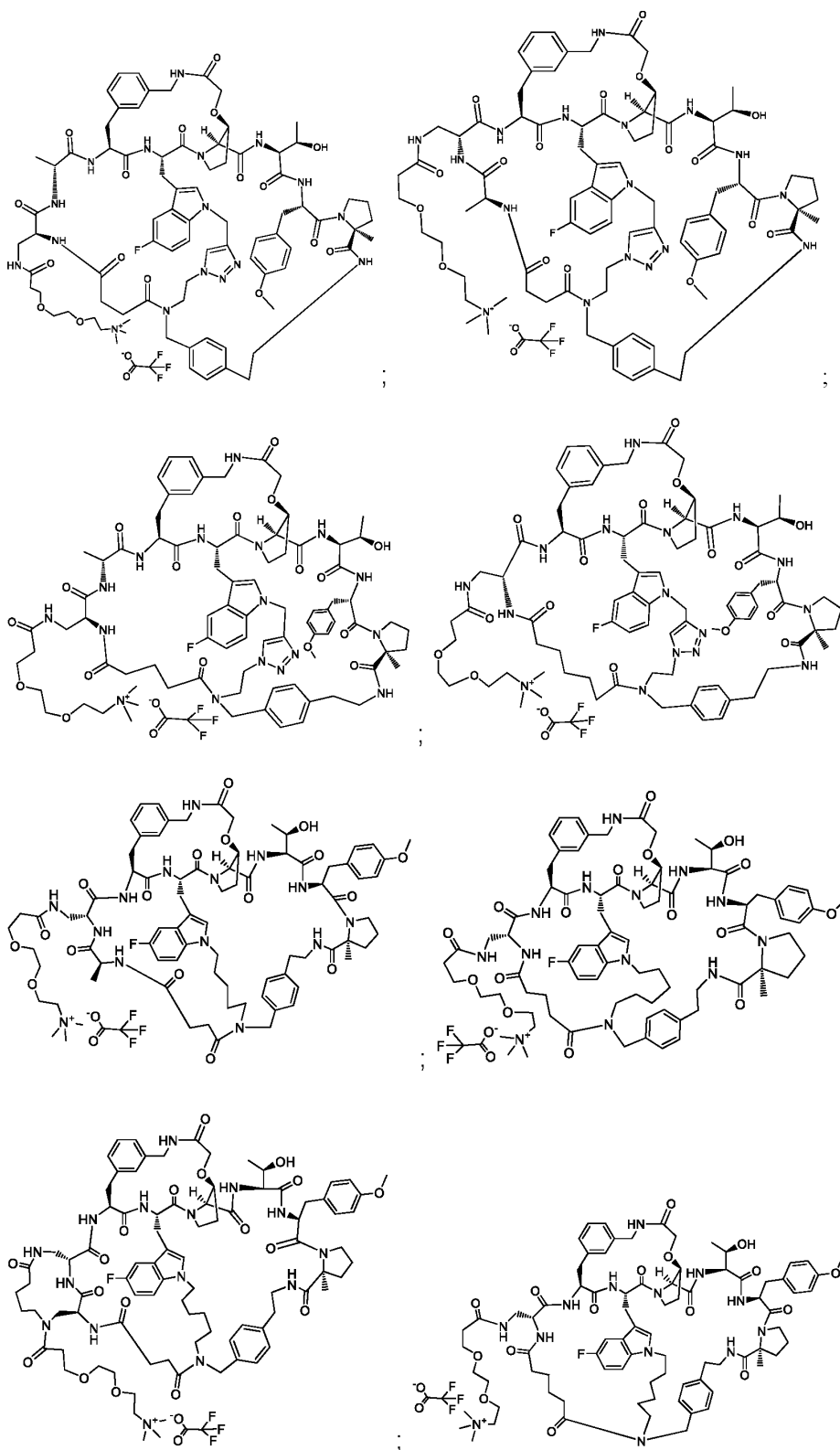


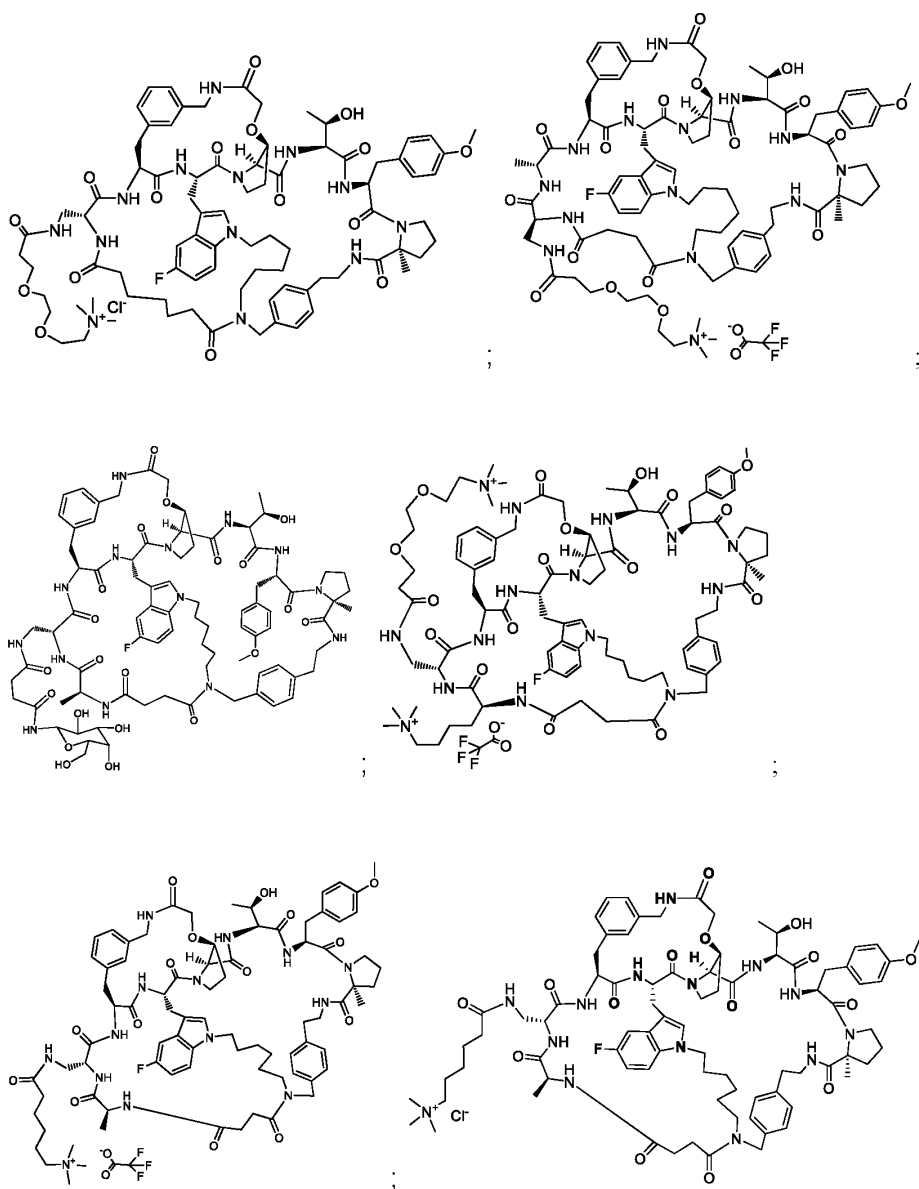
wherein R^{8b} is -H, -CH₃, or -C(CH₃)₃.

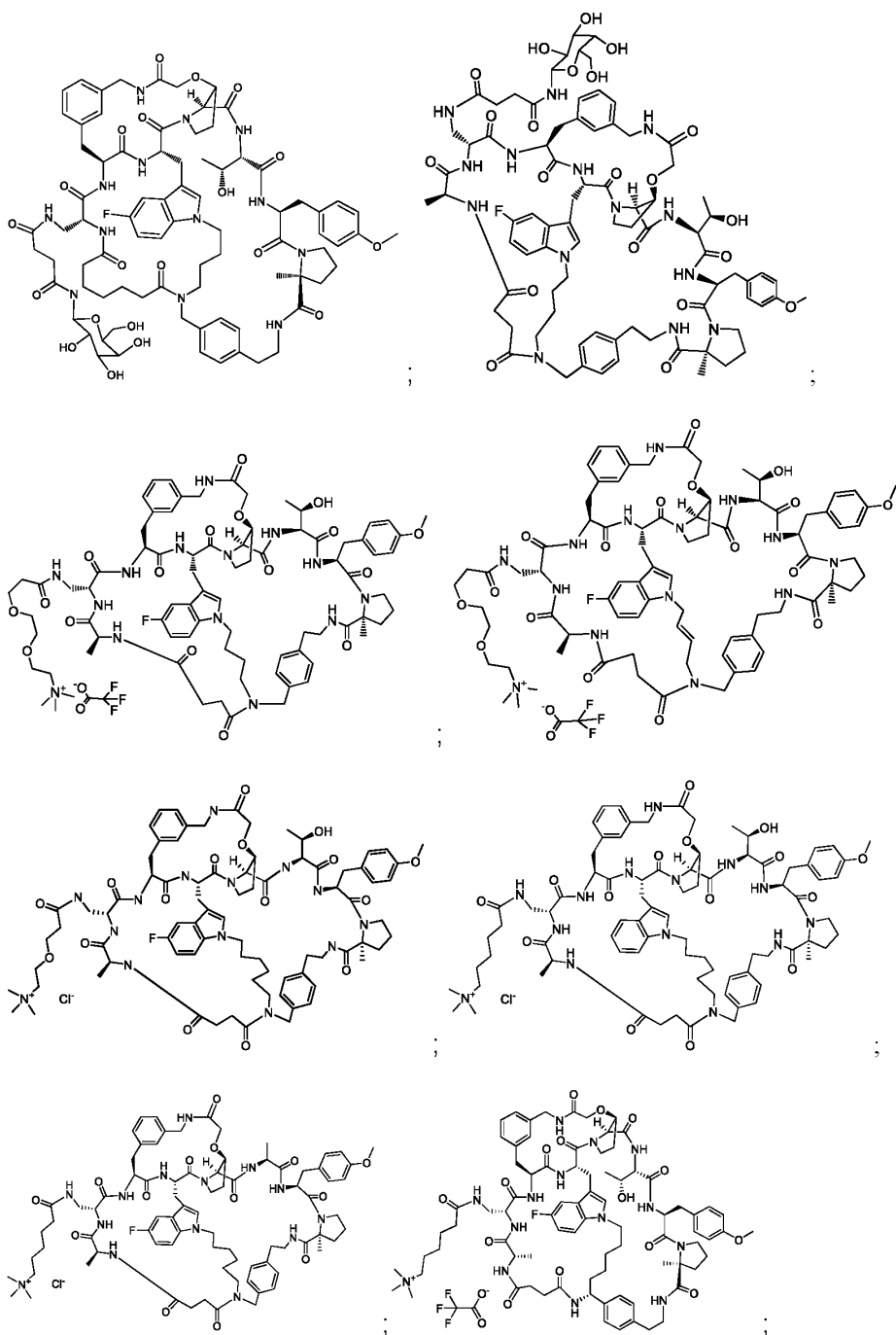
21 A compound of claim 1, which is selected from the group consisting of:

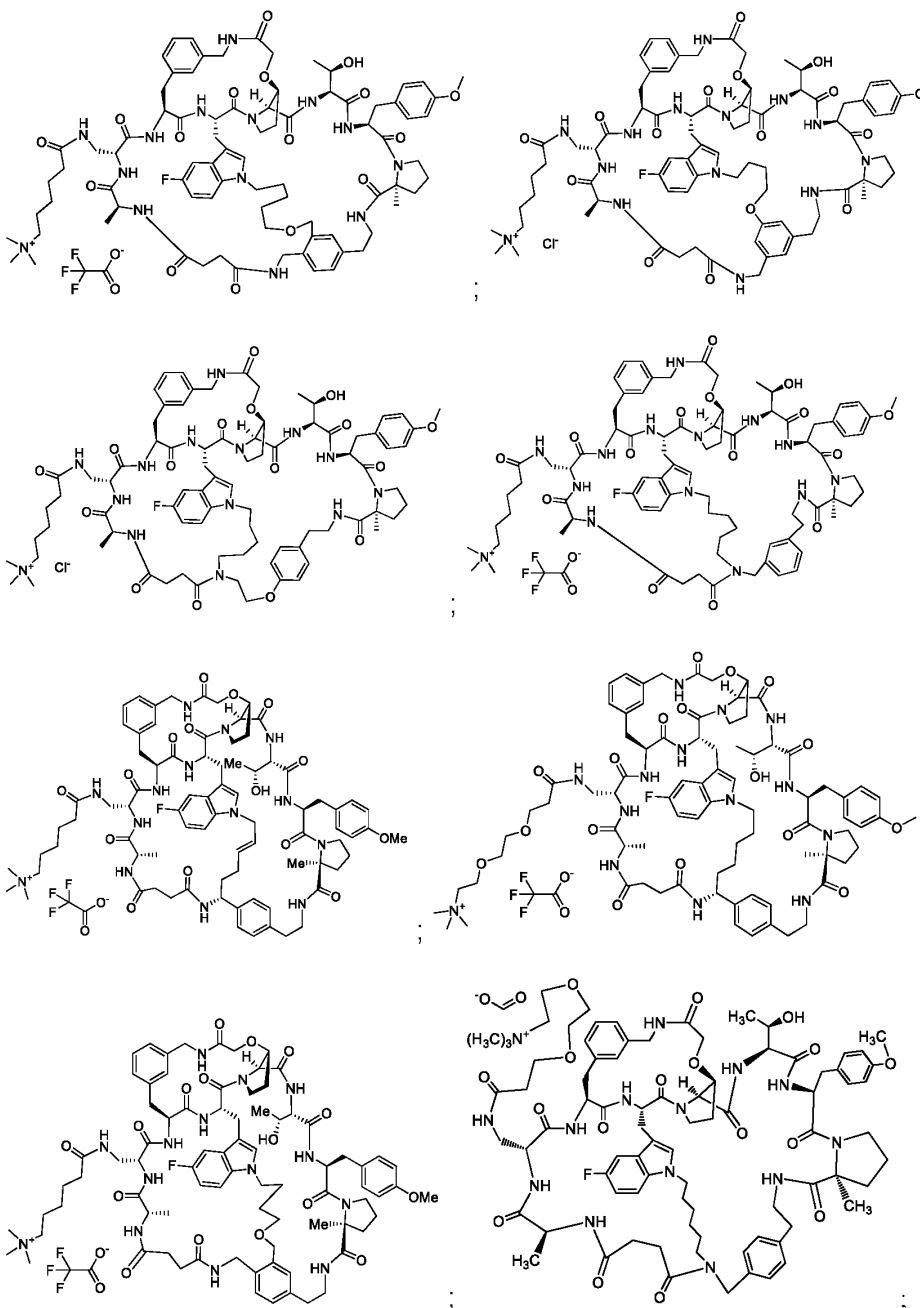


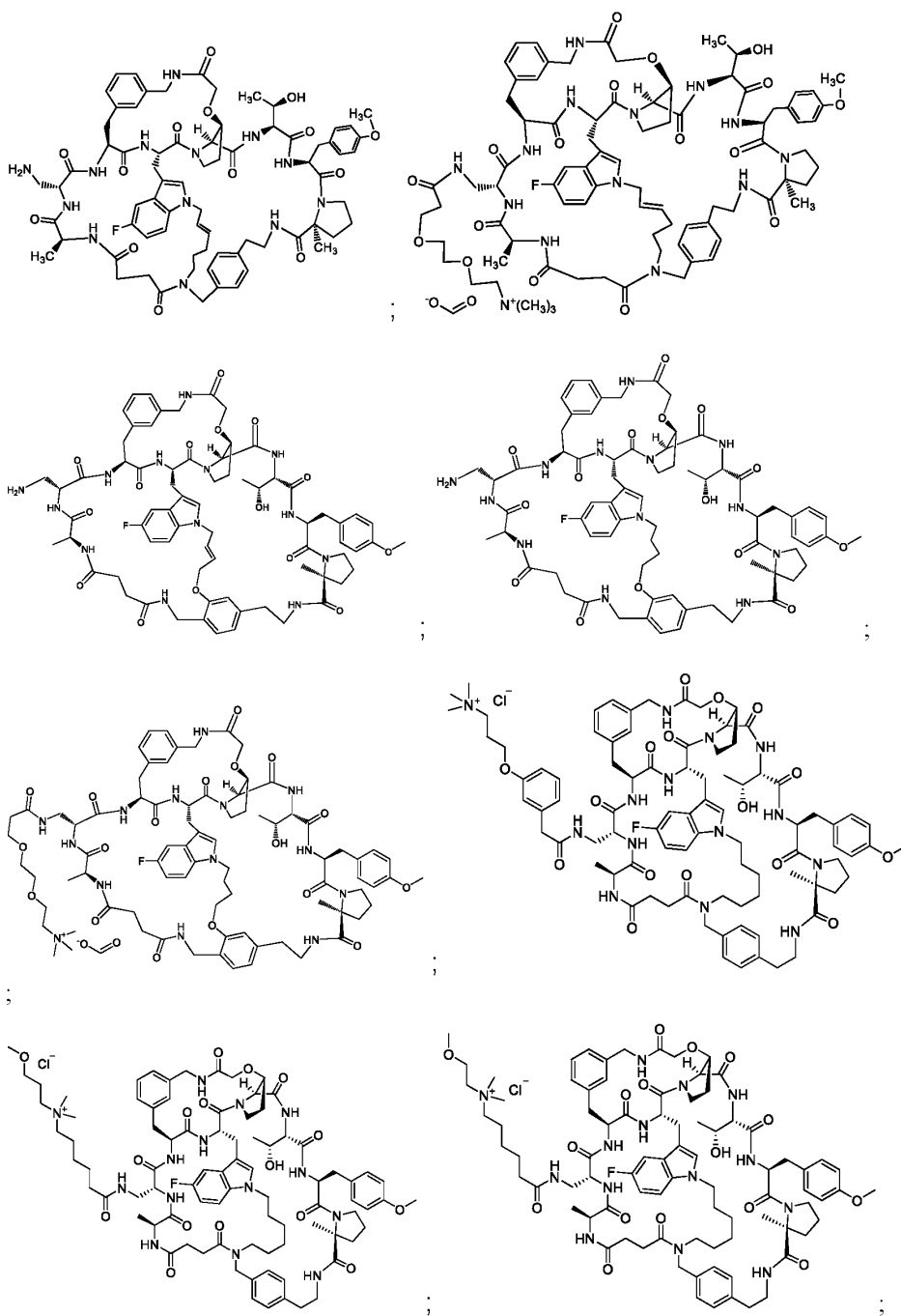


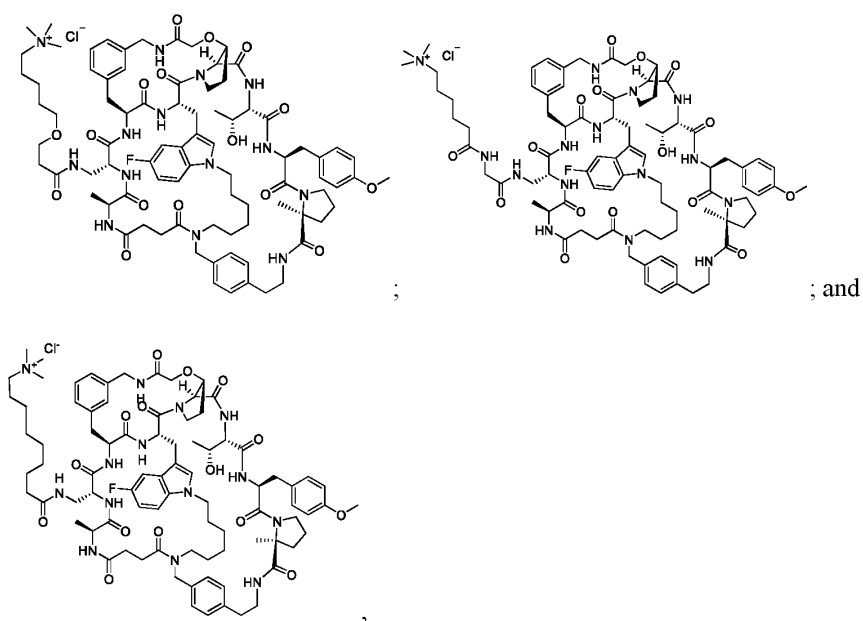












or any pharmaceutically acceptable salt form thereof.

22. A composition comprising at least one compound of any of claims 1 to 21, or a pharmaceutically acceptable salt thereof or freebase form thereof, and at least one pharmaceutically acceptable excipient.
23. A method of treating hypercholesterolemia, comprising administering to a patient in need thereof a therapeutically effective amount of a composition of Claim 22.
24. A method of treating hypercholesterolemia, comprising administering to a patient in need thereof a therapeutically effective amount of a compound of any of Claims 1 to 21.
25. A compound according to any of Claims 1 to 21, or a pharmaceutically acceptable salt thereof, for use in therapy.

26. A compound according to any of Claims 1 to 21, or a pharmaceutically acceptable salt thereof, for treating hypercholesterolemia.